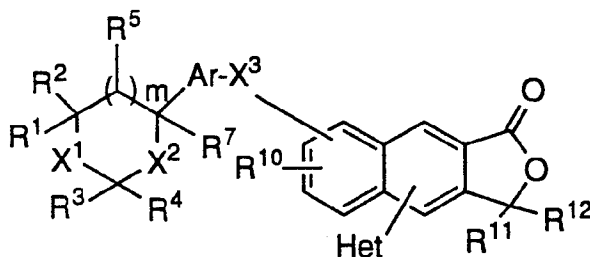




INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

<p>(51) International Patent Classification ⁵ : C07D 407/14, 405/14, 493/08 C07D 409/14, A61K 31/34 A61K 31/38, 31/44 // (C07D 493/08, 311:00, 307:00) (C07D 493/08, 317:00, 311:00)</p>	A1	<p>(11) International Publication Number: WO 94/05658</p> <p>(43) International Publication Date: 17 March 1994 (17.03.94)</p>
<p>(21) International Application Number: PCT/CA93/00339</p> <p>(22) International Filing Date: 19 August 1993 (19.08.93)</p> <p>(30) Priority data: 936,809 27 August 1992 (27.08.92) US</p> <p>(60) Parent Application or Grant (63) Related by Continuation US 936,809 (CON) Filed on 27 August 1992 (27.08.92)</p> <p>(71) Applicant (for all designated States except US): MERCK FROSST CANADA INC. [CA/CA]; 16711 Trans-Canada Highway, Kirkland, Quebec H9H 3L1 (CA).</p> <p>(72) Inventors; and (75) Inventors/Applicants (for US only) : DUCHARME, Yves [CA/CA]; 4501 Kensington, Montreal, Quebec H4B 2W5 (CA). GILLARD, John, W. [AU/CA]; 710 Westchester, Baie d'Urfe, Quebec H9X 2S1 (CA). FORTIN,</p>		<p>Rejean [CA/CA]; 11673 Hurteau, Montreal-Nord, Quebec H1G 2W8 (CA). DUBE, Daniel [CA/CA]; 1811 Blueberry Forest, St. Lazare, Quebec J0P 1V0 (CA). GIRARD, Yves [CA/CA]; 279 Soupras, Ile Bizard, Quebec H3C 2E1 (CA). HAMEL, Pierre [CA/CA]; 1770 Vienne, Vimont, Quebec H7M 2H1 (CA). DELORME, Daniel [CA/CA]; 793 Charbonneau, St. Lazare, Quebec J0P 1V0 (CA).</p> <p>(74) Agents: MURPHY, Kevin, P. et al.; Swabey Ogilvy Renault, Suite 800, 1001 de Maisonneuve Blvd. West, Montreal, Quebec H3A 3C8 (CA).</p> <p>(81) Designated States: AU, BB, BG, BR, BY, CA, CZ, FI, HU, JP, KR, KZ, LK, MG, MN, MW, NO, NZ, PL, RO, RU, SD, SK, UA, US, European patent (AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG).</p> <p>Published With international search report.</p>

(54) Title: HETEROARYLNAPHTHALENE LACTONES AS INHIBITORS OF LEUKOTRIENE BIOSYNTHESIS



(I)

(57) Abstract

Compounds having formula (I): are inhibitors of leukotriene biosynthesis. These compounds are useful as anti-asthmatic, anti-allergic, anti-inflammatory, and cytoprotective agents. They are also useful in treating angina, cerebral spasm, glomerular nephritis, hepatitis, endotoxemia, uveitis, and allograft rejection and in preventing the formation of atherosclerotic plaques.

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TITLE OF THE INVENTION

HETEROARYLNAPHTHALENE LACTONES AS INHIBITORS OF
LEUKOTRIENE BIOSYNTHESIS

5 CROSS-REFERENCE

The hydroxyacid forms of the present lactones are the
subject of a co-pending U.S. patent application, attorney docket no.
18761, U.S. Ser. No. 07/936,810 filed on August 27, 1992, which is
incorporated herein by reference.

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BACKGROUND OF THE INVENTION

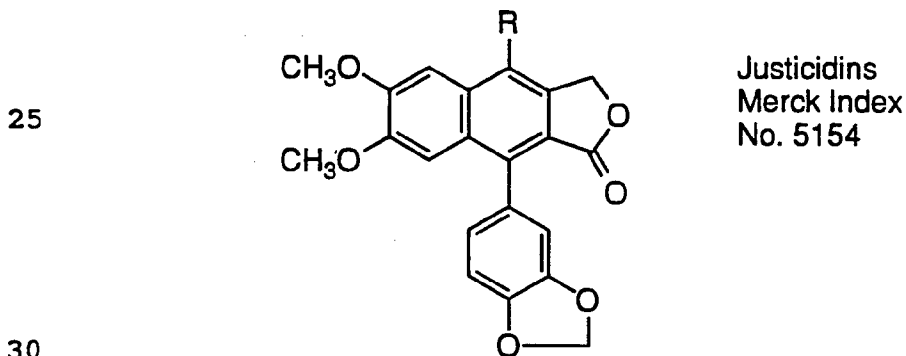
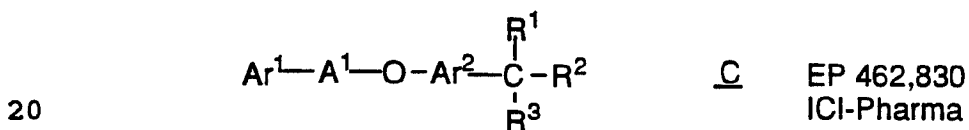
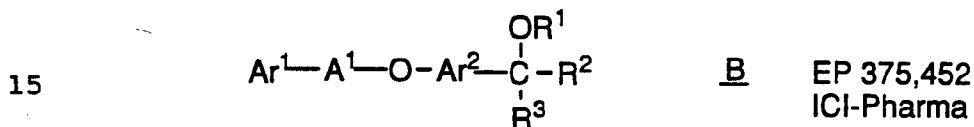
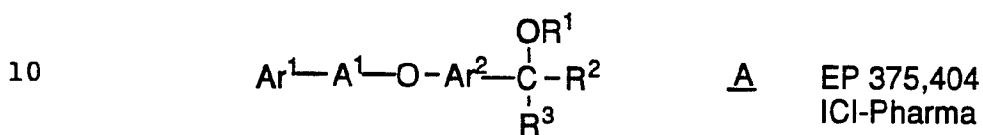
The leukotrienes constitute a group of locally acting
hormones, produced in living systems from arachidonic acid. The
major leukotrienes are Leukotriene B₄ (abbreviated as LTB₄), LTC₄,
15 LTD₄, and LTE₄. The biosynthesis of these leukotrienes begins with
the action of the enzyme 5-lipoxygenase on arachidonic acid to produce
the epoxide known as Leukotriene A₄ (LTA₄), which is converted to
the other leukotrienes by subsequent enzymatic steps. Further details of
the biosynthesis as well as the metabolism of the leukotrienes are to be
20 found in the book Leukotrienes and Lipoxygenases, ed. J. Rokach,
Elsevier, Amsterdam (1989). The actions of the leukotrienes in living
systems and their contribution to various diseases states are also
discussed in the book by Rokach.

European patent application 375,404 (June 27, 1990)
25 describes certain naphthalene containing heterocyclic ethers of structure
A which are inhibitors of the enzyme 5-lipoxygenase. EP application
375,452 (June 27, 1990) describes naphthalene-containing hydrocarbon
ethers of structure B which are reported to possess the same activity.
EP application 462,830 (December 27, 1991) describes bicyclic
30 heterocycle-containing hydrocarbon ethers of structure C which are
reported to possess the same activity. All these series of prior art
compounds differ significantly from the present invention in that they
lack both the aryl substituent and the additional fused ring of the present
compounds.

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A series of natural products known as the justicidins are referred to in the Merck Index, 11th edition, 1989, no. 5154. The justicidins differ considerably from the present compounds in that they are lacking the large pyranylphenyl group.

Hence the compounds of the present invention are completely novel and unexpectedly have biological activity as leukotriene biosynthesis inhibitors.



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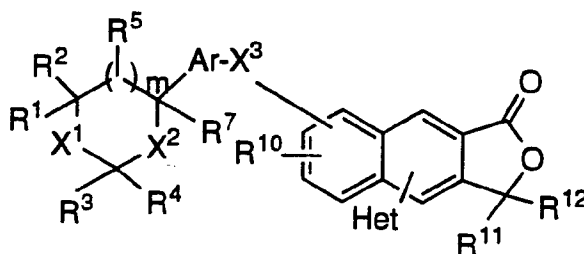
SUMMARY OF THE INVENTION

The present invention relates to heteroarylnaphthalene lactones having activity as leukotriene biosynthesis inhibitors, to methods for their preparation, and to methods and pharmaceutical formulations for using these compounds in mammals (especially humans).

Because of their activity as leukotriene biosynthesis inhibitors, the compounds of the present invention are useful as anti-asthmatic, anti-allergic, anti-inflammatory, and cytoprotective agents. They are also useful in treating angina, cerebral spasm, glomerular nephritis, hepatitis, endotoxemia, uveitis, and allograft rejection and in preventing the formation of atherosclerotic plaques.

DETAILED DESCRIPTION OF THE INVENTION

The compounds of the present invention may be represented by the following formula I:



wherein:

- R1, R5, and R11 is each independently H, OH, lower alkyl, or lower alkoxy;
- R2 is H, lower alkyl, or together with R1 forms a double bonded oxygen (=O);
- R3 is H, lower alkyl, hydroxy lower alkyl, lower alkoxy lower alkyl, or is joined to R1 to form a carbon bridge of 2 or 3 carbon atoms or a mono-oxa carbon bridge of 1 or 2

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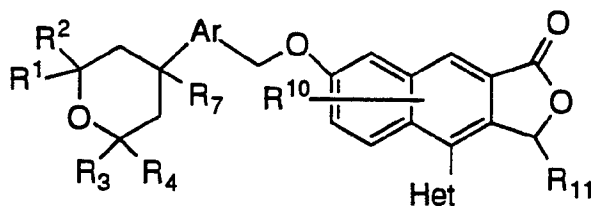
- carbon atoms, said bridge optionally containing a double bond;
- R⁴, R¹², and R¹⁴ is each independently H or lower alkyl;
- 5 R⁶ is H or lower alkyl, or two R⁶ groups on the same or adjacent carbons can form a saturated ring of 3 to 8 members;
- R⁷ is H, OH, lower alkyl, lower alkoxy, lower alkylthio, or lower alkylcarbonyloxy;
- R⁸ is H, halogen, lower alkyl, lower alkoxy, CF₃, CN, COR¹⁴, or a non-bonded electron pair;
- 10 R⁹ and R¹⁰ is each independently H, lower alkyl, lower alkoxy, hydroxy lower alkyl, lower alkoxy lower alkyl, lower alkylthio lower alkyl, lower alkylthio lower alkylcarbonyl, (R⁸)₂-phenylthio lower alkyl, halogen, CN, NO₂, CF₃, N₃, N(R¹³)₂, NR¹³COR¹⁴, NR¹³CON(R¹³)₂, SR¹⁵, S(O)R¹⁵, S(O)₂R¹⁵, S(O)₂N(R¹³)₂, COR¹⁴, CON(R¹³)₂, CO₂R¹⁴, C(R¹⁴)₂OC(R¹⁴)₂-CO₂R¹⁴, or C(R¹⁴)₂CN;
- 15 R¹⁰ is attached to either ring of the naphthalene ring system;
- 20 R¹³ is H or lower alkyl, or two R¹³ groups attached to the same nitrogen may form a saturated ring of 5 or 6 members, optionally containing a second heteroatom chosen from O, S, or NR⁴;
- R¹⁵ is lower alkyl, phenyl-(R⁸)₂, or CF₃;
- 25 X¹ is O, S, S(O), S(O)₂, or C(R⁶)₂;
- X² is O, S, C(R⁶)₂, or bond;
- X³ is C(R⁶)₂S, SC(R⁶)₂, C(R⁶)₂O, OC(R⁶)₂, CR⁶=CR⁶, C(R⁶)₂C(R⁶)₂, O, or S;
- Ar is arylene(R⁸)₂ wherein arylene is a 5-membered aromatic ring containing one O or S and 0-2 nitrogen atoms; a 5-membered aromatic ring containing 1-3 nitrogen atoms; a 6-membered aromatic ring containing 0-3 nitrogen atoms; 2- or 4-pyranone; or 2- or 4-pyridinone;
- 30 Het is heteroaryl(R⁹)₂ wherein heteroaryl is a 5-membered

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aromatic ring containing one O or S and 0-3 nitrogen
atoms; a 5-membered aromatic ring containing 1-4
nitrogen atoms; a 6-membered aromatic ring containing 1-
3 nitrogen atoms; 2- or 4-pyranone; 2- or 4-pyridinone; or
a bicyclic 8-, 9- or 10-membered aromatic ring containing
1-4 heteroatoms chosen independently from O, S, and N.
Het is attached to either ring of the naphthalene ring system; and
m is 0 or 1;
or a pharmaceutically acceptable salt thereof.

The compounds of Formula I are preferably administered
in the hydroxyacid form, which can be prepared by treating the lactones
herein by methods known in the art such as with a strong base.
Discussions herein of dosages, compositions, combinations with other
drugs, etc. can be applied to said hydroxyacid form.

A preferred embodiment of the present invention is
represented by Formula Ia:



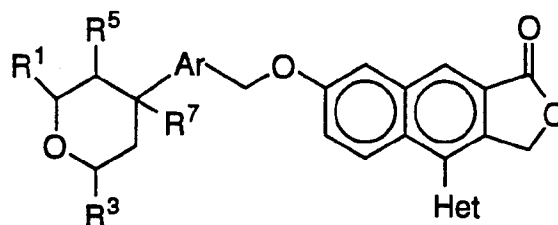
Ia

wherein the substituents are as defined for Formula I or a
pharmaceutically acceptable salt thereof.

Another embodiment of the present invention is
represented by Formula Ib

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5



1b

10

wherein:

R¹ is H, Me, or OMe;R³ is H or Me;R¹ and R³ together are -CH=CH-, -CH₂CH₂-, -OCH₂-, or -CH₂O-;

15

R⁵ is H or OCH₃;R⁷ is OH, OCH₃, or CH₂CH₃;

Ar is Phe; 5,3-Pye; 6,2-Pye; 2,4-Pye; or 4,2-Pye; and

Het is 3-Fu or 3-Th;

or a pharmaceutically acceptable salt thereof.

20

Definitions

The following abbreviations have the indicated meanings:

25	Ac	=	acetyl
	Bn	=	benzyl
	c-Bu	=	cyclobutyl
	Bu ₄ NF	=	tetrabutylammonium fluoride
	c-Pen	=	cyclopentyl
	c-Pr	=	cyclopropyl
30	c-Hex	=	cyclohexyl
	i-Pr	=	isopropyl
	n-Pr	=	normal propyl
	n-Bu	=	normal butyl
	i-Bu	=	isobutyl
	s-Bu	=	secondary butyl

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	t-Bu	=	tertiary butyl
	Et	=	ethyl
	Fu	=	2- or 3-furyl
5	Me	=	methyl
	Ph	=	phenyl
	Py	=	2-, 3- or 4-pyridyl
	Th	=	2- or 3-thienyl
	DHP	=	3,4-dihydro-2H-pyran
10	DMF	=	N,N-dimethylformamide
	DMSO	=	dimethylsulfoxide
	NBS	=	N-bromosuccinimide
	PCC	=	pyridinium chlorochromate
	Phe	=	benzenediyl
15	Pye	=	pyridindiyl
	Super-Hydride	=	lithium triethylborohydride
	TFA	=	trifluoroacetic acid
	THF	=	tetrahydrofuran

20 Alkyl is intended to include linear and branched structures and combinations thereof.

The term "alkyl" includes "lower alkyl" and extends to cover carbon fragments having up to 20 carbon atoms. Examples of alkyl groups include octyl, nonyl, undecyl, dodecyl, tridecyl, tetradecyl, 25 pentadecyl, eicosyl, 3,7-diethyl-2,2-dimethyl-4-propylnonyl, and the like.

"Lower alkyl" means alkyl groups of from 1 to 7 carbon atoms. Examples of lower alkyl groups include methyl, ethyl, propyl, isopropyl, butyl, s- and t-butyl, pentyl, hexyl, heptyl, and the like.

30 "Lower alkoxy" means alkoxy groups of from 1 to 7 carbon atoms of a straight, branched, or cyclic configuration. Examples of lower alkoxy groups include methoxy, ethoxy, propoxy, isopropoxy, cyclopropyloxy, cyclohexyloxy, and the like.

"Lower alkylcarbonyl" means alkylcarbonyl groups of from 1 to 8 carbon atoms of a straight, branched, or cyclic

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configuration. Examples of lower alkylcarbonyl groups are formyl, 2-methylbutanoyl, and cyclohexylacetyl. By way of illustration, the 2-methylbutanoyl group signifies $\text{-COCH(CH}_3\text{)CH}_2\text{CH}_3$.

5 "Hydroxy lower alkyl" means a lower alkyl group carrying a hydroxy group; e.g., $\text{-CH}_2\text{CH(OH)CH}_2\text{CH}_3$.

"Lower alkoxy lower alkyl" means a lower alkyl group carrying a lower alkoxy group; e.g., $\text{-CH}_2\text{CH}_2\text{OCH}_3$.

10 "Lower alkylthio" means alkylthio groups of from 1 to 7 carbon atoms of a straight, branched, or cyclic configuration. Examples of lower alkylthio groups include methylthio, ethylthio, isopropylthio, cyclobutylthio, and the like.

"Lower alkylthio lower alkyl" means a lower alkyl group carrying a lower alkylthio group; e.g., $\text{-CH}_2\text{CH}_2\text{S-c-Pr}$.

15 "Lower alkylthio lower alkylcarbonyl" means a lower alkylcarbonyl group carrying a lower alkylthio group; e.g., $\text{-COCH}_2\text{SCH}_2\text{CH}_3$.

"(R⁸)₂-phenylthio lower alkyl" means a lower alkyl group carrying a phenylthio group which in turn carries two R⁸ substituents; e.g., $\text{-CH}_2\text{CH}_2\text{S-Ph-4-CN}$.

20 Examples of "arylene" are benzene, furan, thiophene, isoxazole, isothiazole, oxazole, thiazole, 1,3,4-oxadiazole, 1,3,4-thiadiazole, 1,2,5-oxadiazole, 1,2,5-thiadiazole, pyrrole, pyrazole, imidazole, 1,3,4-triazole, pyridine, pyrazine, pyrimidine, pyridazine, 1,2,3-triazine, 1,2,4-triazine, 1,3,5-triazine, and the like.

25 Examples of "heteroaryl" are furan, thiophene, oxazole, thiazole, isoxazole, isothiazole, 1,3,4-oxadiazole, 1,3,4-thiadiazole, 1,2,5-oxadiazole, 1,2,5-thiadiazole, pyrrole, pyrazole, imidazole, 1,3,4-triazole, tetrazole, pyridine, pyrazine, pyrimidine, pyridazine, 1,2,3-triazine, 1,2,4-triazine, 1,3,5-triazine, thieno [2,3-b]furan, thieno [3,2-b]pyrrole, indole, benzofuran, benzothiophene, benzimidazole, benzoxazole, benzothiazole, benzo[2,1,3]thiadiazole, thieno[3,2-b]pyridine, furano[2,3-c]pyridine, furano[3,2-b]pyridine, quinoline, isoquinoline, quinoxaline, quinazoline, cinnoline, phthalazine, 1,8-naphthyridine, pteridine, and the like.

30

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Halogen includes F, Cl, Br, and I.

It is intended that the definitions of any substituent (e.g., R⁶, R⁸, etc.) in a particular molecule be independent of its definitions elsewhere in the molecule. Thus, C(R⁶)₂S represents CHCH₃S,
5 C(CH₃)₂S, etc.

Optical Isomers - Diastereomers - Geometric Isomers

Some of the compounds described herein contain one or more asymmetric centers and may thus give rise to diastereomers and
10 optical isomers. The present invention is meant to comprehend such possible diastereomers as well as their racemic and resolved, enantiomerically pure forms, and pharmaceutically acceptable salts thereof.

Some of the compounds described herein contain olefinic
15 double bonds, and unless specified otherwise, are meant to include both E and Z geometric isomers.

Salts

The pharmaceutical compositions of the present invention
20 comprise a compound of Formula I as an active ingredient or a pharmaceutically acceptable salt, thereof, and may also contain a pharmaceutically acceptable carrier and optionally other therapeutic ingredients. The term "pharmaceutically acceptable salts" refers to salts prepared from pharmaceutically acceptable non-toxic bases including
25 inorganic bases and organic bases. Salts derived from inorganic bases include aluminum, ammonium, calcium, copper, ferric, ferrous, lithium, magnesium, manganic salts, manganous, potassium, sodium, zinc and the like. Particularly preferred are the ammonium, calcium, magnesium, potassium and sodium salts. Salts derived from
30 pharmaceutically acceptable organic non-toxic bases include salts of primary, secondary, and tertiary amines, substituted amines including naturally occurring substituted amines, cyclic amines and basic ion exchange resins, such as arginine, betaine, caffeine, choline, N,N'-dibenzylethylenediamine, diethylamine, 2-diethylaminoethanol, 2-

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dimethylaminoethanol, ethanolamine, ethylenediamine, N-ethylmorpholine, N-ethylpiperidine, glucamine, glucosamine, histidine, hydrabamine, isopropylamine, lysine, methylglucamine, morpholine, piperazine, piperidine, polyamine resins, procaine, purines, theobromine, triethylamine, trimethylamine, tripropylamine, tromethamine, and the like.

When the compound of the present invention is basic, salts may be prepared from pharmaceutically acceptable non-toxic acids, including inorganic and organic acids. Such acids include acetic, benzenesulfonic, benzoic, camphorsulfonic, citric, ethanesulfonic, fumaric, gluconic, glutamic, hydrobromic, hydrochloric, isethionic, lactic, maleic, malic, mandelic, methanesulfonic, mucic, nitric, pantoic, pantothenic, phosphoric, succinic, sulfuric, tartaric, p-toluenesulfonic acid, and the like. Particularly preferred are citric, hydrobromic, hydrochloric, maleic, phosphoric, sulfuric, and tartaric acids.

It will be understood that in the discussion of methods of treatment which follows, references to the compounds of Formula I are meant to also include the pharmaceutically acceptable salts.

Utilities

The ability of the compounds of Formula I to inhibit biosynthesis of the leukotrienes makes them useful for preventing or reversing the symptoms induced by the leukotrienes in a human subject. This inhibition of the mammalian biosynthesis of leukotrienes indicates that the compounds and pharmaceutical compositions thereof are useful to treat, prevent, or ameliorate in mammals and especially in humans: 1) pulmonary disorders including diseases such as asthma, chronic bronchitis, and related obstructive airway diseases, 2) allergies and allergic reactions such as allergic rhinitis, contact dermatitis, allergic conjunctivitis, and the like, 3) inflammation such as arthritis or inflammatory bowel disease, 4) pain, 5) skin disorders such as psoriasis, atopic eczema, and the like, 6) cardiovascular disorders such as angina, formation of atherosclerotic plaques, myocardial ischemia, hypertension, platelet aggregation, and the like, 7) renal insufficiency

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arising from ischaemia induced by immunological or chemical (cyclosporin) etiology, 8) migraine or cluster headache, 9) ocular conditions such as uveitis, 10) hepatitis resulting from chemical, immunological or infectious stimuli, 11) trauma or shock states such as burn injuries, endotoxemia, and the like, 12) allograft rejection, 13) prevention of side effects associated with therapeutic administration of cytokines such as Interleukin II and tumor necrosis factor, 14) chronic lung diseases such as cystic fibrosis, bronchitis and other small- and large-airway diseases, 15) cholecystitis, and 16) multiple sclerosis.

Thus, the compounds of the present invention may also be used to treat or prevent mammalian (especially, human) disease states such as erosive gastritis; erosive esophagitis; diarrhea; cerebral spasm; premature labor; spontaneous abortion; dysmenorrhea; ischemia; noxious agent-induced damage or necrosis of hepatic, pancreatic, renal, or myocardial tissue; liver parenchymal damage caused by hepatotoxic agents such as CCl₄ and D-galactosamine; ischemic renal failure; disease-induced hepatic damage; bile salt induced pancreatic or gastric damage; trauma- or stress-induced cell damage; and glycerol-induced renal failure. The compounds also act as inhibitors of tumor metastasis and exhibit cytoprotective action.

The cytoprotective activity of a compound may be observed in both animals and man by noting the increased resistance of the gastrointestinal mucosa to the noxious effects of strong irritants, for example, the ulcerogenic effects of aspirin or indomethacin. In addition to lessening the effect of non-steroidal anti-inflammatory drugs on the gastrointestinal tract, animal studies show that cytoprotective compounds will prevent gastric lesions induced by oral administration of strong acids, strong bases, ethanol, hypertonic saline solutions, and the like.

Two assays can be used to measure cytoprotective ability. These assays are: (A) an ethanol-induced lesion assay, and (B) an indomethacin-induced ulcer assay and are described in EP 140,684.

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Dose Ranges

5 The magnitude of prophylactic or therapeutic dose of a compound of Formula I will, of course, vary with the nature of the severity of the condition to be treated and with the particular compound of Formula I and its route of administration. It will also vary according to the age, weight and response of the individual patient. In general, the daily dose range for anti-asthmatic, anti-allergic or anti-inflammatory use and generally, uses other than cytoprotection, lie within the range of from about 0.001 mg to about 100 mg per kg body weight of a mammal, preferably 0.01 mg to about 10 mg per kg, and most preferably 0.1 to 1 mg per kg, in single or divided doses. On the other hand, it may be necessary to use dosages outside these limits in some cases.

15 For use where a composition for intravenous administration is employed, a suitable dosage range for anti-asthmatic, anti-inflammatory or anti-allergic use is from about 0.001 mg to about 25 mg (preferably from 0.01 mg to about 1 mg) of a compound of Formula I per kg of body weight per day and for cytoprotective use from about 0.1 mg to about 100 mg (preferably from about 1 mg to about 100 mg and more preferably from about 1 mg to about 10 mg) of a compound of Formula I per kg of body weight per day.

25 In the case where an oral composition is employed, a suitable dosage range for anti-asthmatic, anti-inflammatory or anti-allergic use is, e.g. from about 0.01 mg to about 100 mg of a compound of Formula I per kg of body weight per day, preferably from about 0.1 mg to about 10 mg per kg and for cytoprotective use from 0.1 mg to about 100 mg (preferably from about 1 mg to about 100 mg and more preferably from about 10 mg to about 100 mg) of a compound of Formula I per kg of body weight per day.

30 For the treatment of diseases of the eye, ophthalmic preparations for ocular administration comprising 0.001-1% by weight solutions or suspensions of the compounds of Formula I in an acceptable ophthalmic formulation may be used.

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5 The exact amount of a compound of the Formula I to be used as a cytoprotective agent will depend on, inter alia, whether it is being administered to heal damaged cells or to avoid future damage, on the nature of the damaged cells (e.g., gastrointestinal ulcerations vs. nephrotic necrosis), and on the nature of the causative agent. An example of the use of a compound of the Formula I in avoiding future damage would be co-administration of a compound of the Formula I with a non-steroidal anti-inflammatory drug that might otherwise cause such damage (for example, indomethacin). For such use, the compound of Formula I is administered from 30 minutes prior up to 30 minutes after administration of the NSAID. Preferably it is administered prior to or simultaneously with the NSAID, (for example, in a combination dosage form).

15 Pharmaceutical Compositions

Any suitable route of administration may be employed for providing a mammal, especially a human with an effective dosage of a compound of the present invention. For example, oral, rectal, topical, parenteral, ocular, pulmonary, nasal, and the like may be employed. Dosage forms include tablets, troches, dispersions, suspensions, solutions, capsules, creams, ointments, aerosols, and the like.

25 The pharmaceutical compositions of the present invention comprise a compound of Formula I as an active ingredient or a pharmaceutically acceptable salt thereof, and may also contain a pharmaceutically acceptable carrier and optionally other therapeutic ingredients. The term "pharmaceutically acceptable salts" refers to salts prepared from pharmaceutically acceptable non-toxic bases or acids including inorganic bases or acids and organic bases or acids.

30 The compositions include compositions suitable for oral, rectal, topical, parenteral (including subcutaneous, intramuscular, and intravenous), ocular (ophthalmic), pulmonary (nasal or buccal inhalation), or nasal administration, although the most suitable route in any given case will depend on the nature and severity of the conditions being treated and on the nature of the active ingredient. They may be

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conveniently presented in unit dosage form and prepared by any of the methods well-known in the art of pharmacy.

5 For administration by inhalation, the compounds of the present invention are conveniently delivered in the form of an aerosol spray presentation from pressurized packs or nebulisers. The compounds may also be delivered as powders which may be formulated and the powder composition may be inhaled with the aid of an insufflation powder inhaler device. The preferred delivery system for inhalation is a metered dose inhalation (MDI) aerosol, which may be
10 formulated as a suspension or solution of compound I in suitable propellants, such as fluorocarbons or hydrocarbons.

Suitable topical formulations of Compound I include transdermal devices, aerosols, creams, ointments, lotions, dusting
15 powders, and the like.

In practical use, the compounds of Formula I can be combined as the active ingredient in intimate admixture with a pharmaceutical carrier according to conventional pharmaceutical compounding techniques. The carrier may take a wide variety of forms depending on the form of preparation desired for administration, e.g.,
20 oral or parenteral (including intravenous). In preparing the compositions for oral dosage form, any of the usual pharmaceutical media may be employed, such as, for example, water, glycols, oils, alcohols, flavoring agents, preservatives, coloring agents and the like in the case of oral liquid preparations, such as, for example, suspensions,
25 elixirs and solutions; or carriers such as starches, sugars, microcrystalline cellulose, diluents, granulating agents, lubricants, binders, disintegrating agents and the like in the case of oral solid preparations such as, for example, powders, capsules and tablets, with the solid oral preparations being preferred over the liquid preparations.
30 Because of their ease of administration, tablets and capsules represent the most advantageous oral dosage unit form in which case solid pharmaceutical carriers are obviously employed. If desired, tablets may be coated by standard aqueous or nonaqueous techniques.

- 15 -

In addition to the common dosage forms set out above, the compounds of Formula I may also be administered by controlled release means and/or delivery devices such as those described in U.S. Patent Nos. 3,845,770; 3,916,899; 3,536,809; 3,598,123; 3,630,200 and
5 4,008,719, the disclosures of which are hereby incorporated herein by reference.

Pharmaceutical compositions of the present invention suitable for oral administration may be presented as discrete units such as capsules, cachets or tablets each containing a predetermined amount
10 of the active ingredient, as a powder or granules or as a solution or a suspension in an aqueous liquid, a non-aqueous liquid, an oil-in-water emulsion or a water-in-oil liquid emulsion. Such compositions may be prepared by any of the methods of pharmacy but all methods include the step of bringing into association the active ingredient with the carrier
15 which constitutes one or more necessary ingredients. In general, the compositions are prepared by uniformly and intimately admixing the active ingredient with liquid carriers or finely divided solid carriers or both, and then, if necessary, shaping the product into the desired presentation. For example, a tablet may be prepared by compression or
20 molding, optionally with one or more accessory ingredients. Compressed tablets may be prepared by compressing in a suitable machine, the active ingredient in a free-flowing form such as powder or granules, optionally mixed with a binder, lubricant, inert diluent, surface active or dispersing agent. Molded tablets may be made by
25 molding in a suitable machine, a mixture of the powdered compound moistened with an inert liquid diluent. Desirably, each tablet contains from about 2.5 mg to about 500 mg of the active ingredient and each cachet or capsule contains from about 2.5 to about 500 mg of the active
30 ingredient.

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The following are examples of representative pharmaceutical dosage forms for the compounds of Formula I:

5	<u>Injectable Suspension (I.M.)</u>	<u>mg/mL</u>
	Compound of Formula I	10
	Methylcellulose	5.0
	Tween 80	0.5
	Benzyl alcohol	9.0
10	Benzalkonium chloride	1.0
	Water for injection to a total volume of 1 mL	
15	<u>Tablet</u>	<u>mg/tablet</u>
	Compound of Formula I	25
	Microcrystalline Cellulose	415
	Povidone	14.0
	Pregelatinized Starch	43.5
	Magnesium Stearate	2.5
		<hr/> 500
20	<u>Capsule</u>	<u>mg/capsule</u>
	Compound of Formula I	25
	Lactose Powder	573.5
	Magnesium Stearate	1.5
25		<hr/> 600
30	<u>Aerosol</u>	<u>Per canister</u>
	Compound of Formula I	24 mg
	Lecithin, NF Liquid Concentrate	1.2 mg
	Trichlorofluoromethane, NF	4.025 g
	Dichlorodifluoromethane, NF	12.15 g

Combinations with other drugs

In addition to the compounds of Formula I, the pharmaceutical compositions of the present invention can also contain

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other active ingredients, such as cyclooxygenase inhibitors, non-steroidal anti-inflammatory drugs (NSAIDs), peripheral analgesic agents such as zomepirac diflunisal and the like. The weight ratio of the compound of the Formula I to the second active ingredient may be varied and will depend upon the effective dose of each ingredient. Generally, an effective dose of each will be used. Thus, for example, when a compound of the Formula I is combined with an NSAID the weight ratio of the compound of the Formula I to the NSAID will generally range from about 1000:1 to about 1:1000, preferably about 200:1 to about 1:200. Combinations of a compound of the Formula I and other active ingredients will generally also be within the aforementioned range, but in each case, an effective dose of each active ingredient should be used.

NSAIDs can be characterized into five groups:

- (1) the propionic acid derivatives;
- (2) the acetic acid derivatives;
- (3) the fenamic acid derivatives;
- (4) the oxicams; and
- (5) the biphenylcarboxylic acid derivatives;

or a pharmaceutically acceptable salt thereof.

The propionic acid derivatives which may be used comprise: alminoprofen, benoxaprofen, bucloxic acid, carprofen, fenbufen, fenoprofen, fluprofen, flurbiprofen, ibuprofen, indoprofen, ketoprofen, miroprofen, naproxen, oxaprozin, pirprofen, prano-profen, suprofen, tiaprofenic acid, and tioxaprofen. Structurally related propionic acid derivatives having similar analgesic and anti-inflammatory properties are also intended to be included in this group.

Thus, "propionic acid derivatives" as defined herein are non-narcotic analgesics/non-steroidal anti-inflammatory drugs having a free $-\text{CH}(\text{CH}_3)\text{COOH}$ or $-\text{CH}_2\text{CH}_2\text{COOH}$ group (which optionally can be in the form of a pharmaceutically acceptable salt group, e.g., $-\text{CH}(\text{CH}_3)\text{COO}-\text{Na}^+$ or $-\text{CH}_2\text{CH}_2\text{COO}-\text{Na}^+$), typically attached directly or via a carbonyl function to a ring system, preferably to an aromatic ring system.

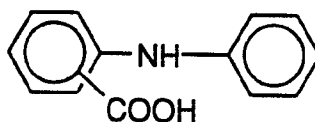
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The acetic acid derivatives which may be used comprise: indomethacin, which is a preferred NSAID, acemetacin, alclofenac, clidanac, diclofenac, fenclofenac, fenclozic acid, fentiazac, furofenac, ibufenac, isoxepac, oxpinac, sulindac, tiopinac, tolmetin, zidometacin and zomepirac. Structurally related acetic acid derivatives having similar analgesic and anti-inflammatory properties are also intended to be encompassed by this group.

Thus, "acetic acid derivatives" as defined herein are non-narcotic analgesics/non-steroidal anti-inflammatory drugs having a free -CH₂COOH group (which optionally can be in the form of a pharmaceutically acceptable salt group, e.g. -CH₂COO-Na⁺), typically attached directly to a ring system, preferably to an aromatic or heteroaromatic ring system.

The fenamic acid derivatives which may be used comprise: flufenamic acid, meclofenamic acid, mefenamic acid, niflumic acid and tolfenamic acid. Structurally related fenamic acid derivatives having similar analgesic and anti-inflammatory properties are also intended to be encompassed by this group.

Thus, "fenamic acid derivatives" as defined herein are non-narcotic analgesics/non-steroidal anti-inflammatory drugs which contain the basic structure:



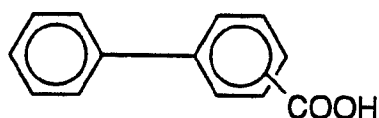
which can bear a variety of substituents and in which the free -COOH group can be in the form of a pharmaceutically acceptable salt group, e.g., -COO-Na⁺.

The biphenylcarboxylic acid derivatives which can be used comprise: diflunisal and flufenisal. Structurally related biphenylcarboxylic acid derivatives having similar analgesic and anti-inflammatory properties are also intended to be encompassed by this group.

- 19 -

Thus, "biphenylcarboxylic acid derivatives" as defined herein are non-narcotic analgesics/non-steroidal anti-inflammatory drugs which contain the basic structure:

5

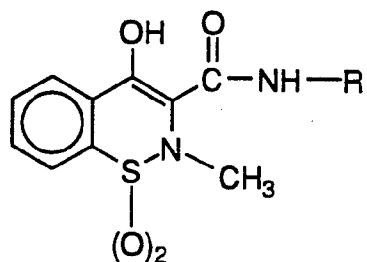


which can bear a variety of substituents and in which the free -COOH group can be in the form of a pharmaceutically acceptable salt group, e.g., -COO-Na⁺.

The oxicams which can be used in the present invention comprise: isoxicam, piroxicam, sudoxicam and tenoxicam. Structurally related oxicams having similar analgesic and anti-inflammatory properties are also intended to be encompassed by this group.

Thus, "oxicams" as defined herein are non-narcotic analgesics/non-steroidal anti-inflammatory drugs which have the general formula:

20



25

wherein R is an aryl or heteroaryl ring system.

The following NSAIDs may also be used: amfenac sodium, aminoprofen, anitrazafen, antrafenine, auranofin, bendazac lysinate, benzydanine, beprozin, broperamole, bufezolac, cinmetacin, ciproquazone, cloximate, dazidamine, deboxamet, delmetacin, detomidine, dexindoprofen, diacerein, di-fisalamine, difenpyramide, emorfazone, enfenamic acid, enolicam, epirizole, etersalate, etodolac, etofenamate, fanetizole mesylate, fenclorac, fendosal, fenflumizole, feprazone, floctafenine, flunixin, flunoxaprofen, fluproquazone,

- 20 -

fopirtoline, fosfosal, furclopofen, glucametacin, guaimesal, ibuproxam, isofezolac, isonixim, isoprofen, isoxicam, lefetamine HCl, leflunomide, lofemizole, lonazolac calcium, lotifazole, loxoprofen, lysin clonixinate, meclofenamate sodium, meseclazone, nabumetone, nictindole, nimesulide, orpanoxin, oxametacin, oxapadol, perisoxal citrate, pimeprofen, pimetacin, piroxan, pirazolac, pirfenidone, proglumetacin maleate, proquazone, pyridoxiprofen, sudoxicam, talmetacin, talniflumate, tenoxicam, thiazolinobutazone, thielavin B, tiaramide HCl, tifihamizole, timegadine, tolpadol, tryptamid, and ufenamate.

The following NSAIDs, designated by company code number (see e.g., Pharmaprojects), may also be used:

480156S, AA861, AD1590, AFP802, AFP860, AI77B, AP504, AU8001, BPPC, BW540C, CHINOIN 127, CN100, EB382, EL508, F1044, GV3658, ITF182, KCNTEI6090, KME4, LA2851, MR714, MR897, MY309, ONO3144, PR823, PV102, PV108, R830, RS2131, SCR152, SH440, SIR133, SPAS510, SQ27239, ST281, SY6001, TA60, TAI-901 (4-benzoyl-1-indancarboxylic acid), TVX2706, U60257, UR2301, and WY41770.

Finally, NSAIDs which may also be used include the salicylates, specifically acetyl salicylic acid and the phenylbutazones, and pharmaceutically acceptable salts thereof.

In addition to indomethacin, other preferred NSAIDs are acetyl salicylic acid, diclofenac, fenbufen, fenoprofen, flurbiprofen, ibuprofen, ketoprofen, naproxen, phenylbutazone, piroxicam, sulindac, and tolmetin.

Pharmaceutical compositions comprising the Formula I compounds may also contain inhibitors of the biosynthesis of the leukotrienes such as are disclosed in EP 138,481 (April 24, 1985), EP 115,394 (August 8, 1984), EP 136,893 (April 10, 1985), and EP 140,709 (May 8, 1985), which are hereby incorporated herein by reference.

The compounds of the Formula I may also be used in combination with leukotriene antagonists such as those disclosed in EP 106,565 (April 25, 1984) and EP 104,885 (April 4, 1984) which are

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hereby incorporated herein by reference and others known in the art such as those disclosed in EP Application Nos. 56,172 (July 21, 1982) and 61,800 (June 10, 1982); and in U.K. Patent Specification No. 2,058,785 (April 15, 1981), which are hereby incorporated herein by
5 reference.

Pharmaceutical compositions comprising the Formula I compounds may also contain as the second active ingredient, prostaglandin antagonists such as those disclosed in EP 11,067 (May 28, 1980) or thromboxane antagonists such as those disclosed in U.S. Pat.
10 4,237,160. They may also contain histidine decarboxylase inhibitors such as α -fluoromethyl-histidine, described in U.S. Pat. 4,325,961. The compounds of the Formula I may also be advantageously combined with an H₁- or H₂-receptor antagonist, such as for instance acetamazole, aminothiadiazoles disclosed in EP 40,696 (December 2, 1981),
15 benadryl, cimetidine, famotidine, framamine, histadyl, phenergan, ranitidine, terfenadine and like compounds, such as those disclosed in U.S. Patent Nos. 4,283,408; 4,362,736; and 4,394,508. The pharmaceutical compositions may also contain a K⁺/H⁺ ATPase inhibitor such as omeprazole, disclosed in U.S. Pat. 4,255,431, and the
20 like. Compounds of Formula I may also be usefully combined with most cell stabilizing agents, such as 1,3-bis(2-carboxy-chromon-5-yl-oxy)-2-hydroxypropane and related compounds described in British Patent Specifications 1,144,905 and 1,144,906. Another useful pharmaceutical composition comprises the Formula I compounds in
25 combination with serotonin antagonists such as methysergide, the serotonin antagonists described in Nature, Vol. 316, pages 126-131, 1985, and the like. Each of the references referred to in this paragraph is hereby incorporated herein by reference.

Other advantageous pharmaceutical compositions comprise
30 the Formula I compounds in combination with anti-cholinergics such as ipratropium bromide, bronchodilators such as the beta agonist salbutamol, metaproterenol, terbutaline, fenoterol and the like, and the anti-asthmatic drugs theophylline, choline theophyllinate and enprofylline, the calcium antagonists nifedipine, diltiazem, nitrendipine,

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verapamil, nimodipine, felodipine, etc. and the corticosteroids, hydrocortisone, methylprednisolone, betamethasone, dexamethasone, beclomethasone, and the like.

5 Methods of Synthesis

Compounds of the present invention can be prepared according to the following methods. Temperatures are in degrees Celsius. The substituents are the same as in Formula I except where defined otherwise.

10

Scheme I

The route which is used to prepare the lactone VI is outlined in Scheme I. The heteroaryl carboxaldehyde II is converted to the thioacetal III by treatment with thiophenol in the presence of a
15 Lewis-Acid such as $\text{BF}_3 \cdot \text{Et}_2\text{O}$ in an organic solvent such as isopropyl acetate. The thioacetal III is then converted to the lactone V by treatment with a base such as $n\text{-BuLi}$ in an organic solvent such as THF followed by the successive additions of 2-(5H) furanone and the benzyloxy benzaldehyde IV in an organic solvent such as THF followed
20 by quenching with an acid such as HOAc. The cyclization, dehydration and debenzoylation are achieved simultaneously by heating the lactone V in the presence of an acid such as TFA, and using thioanisole as organic solvent providing the lactone VI.

25 Scheme II

The preparation of compounds of Formula I (wherein $\text{X}^3 = -\text{C}(\text{R}^6)_2\text{O}-$) is described in Scheme II. A first method requires coupling of the naphthol VI with a benzylic halide or activated alcohol of type VII (wherein $\text{X} = \text{Cl}, \text{Br}, \text{I}, \text{OMs}, \text{OTs}$) in a polar organic solvent such
30 DMF in the presence of an inorganic base such as Cs_2CO_3 .

In an alternate procedure, the naphthol VI is condensed with the benzylic alcohol VII (wherein $\text{X} = \text{OH}$) in the presence of a phosphine such as Ph_3P and an azodicarboxylate diester, in a solvent

- 23 -

such as THF, to afford Formula I (wherein $X^3 = -C(R^6)_2O-$) compounds.

Scheme III

5 The synthesis of compounds of Formula I (wherein $X^3 = -OCH_2-$) is described in Scheme III. The phenol lactone VI may be converted to the triflate VIII by treatment with trifluoromethane-sulfonic anhydride in the presence of an organic base such as pyridine in
10 a solvent such as CH_2Cl_2 . Subsequent treatment of VIII in a solvent such as DMSO/MeOH with an organic base such as triethylamine, a phosphine such as 1,1'-bis(diphenylphosphino)-ferrocene, a palladium(II) salt such as palladium(II)acetate under an atmosphere of carbon monoxide will lead to the ester IX. The hydrolysis of the ester
15 IX may be achieved using an inorganic base such as lithium hydroxide in water and the resulting acid may be reduced to the alcohol XI by treatment with a chloroformate such as isopropyl chloroformate in the presence of an organic base such as triethylamine in an organic solvent such as THF, followed by addition of a reducing agent such as sodium borohydride in water. The alcohol XI may be then converted to the
20 halide XII by treatment with triphenylphosphine, imidazole and CBr_4 in an organic solvent such as CH_2Cl_2 . Coupling of halide XII with the appropriate phenol XIII in an organic solvent such as DMF using an inorganic base such as K_2CO_3 provides compounds of formula I (wherein $X^3 = -OCH_2-$) of the present invention.
25

Scheme IV

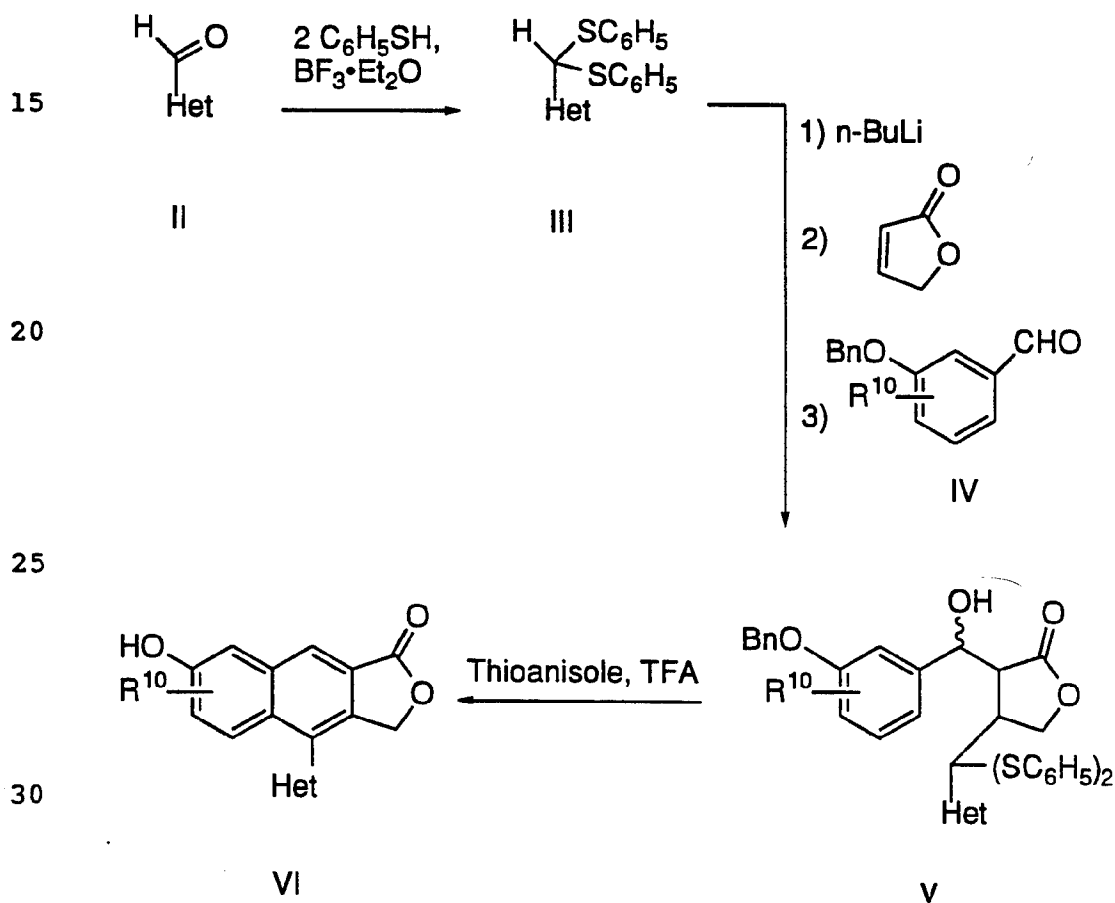
30 Scheme IV illustrates the conversion of compounds of formula I (wherein $R^{11}/R^{12} = H$) to formula I (wherein R^{11}/R^{12} is not H). Compounds of formula I (wherein $R^{11}/R^{12} = H$) may be alkylated by treatment with a strong base such as lithium diisopropylamide in an organic solvent such as THF, followed by quenching with an alkyl halide such as methyl iodide. A second alkylation may be achieved by using the same conditions. Conversion of compounds of formula I (wherein $R^{11}/R^{12} = H$) to compounds I (wherein $R^{11} = OH$, $R^{12} = H$)

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may be achieved by an hydrolysis process using an inorganic base such as NaOH in a solvent such as EtOH/H₂O to provide the carboxylate salt XIV, followed by an oxidation step using an oxidizing agent such as pyridinium chlorochromate in an organic solvent such as CH₂Cl₂. The reaction of compound I (R¹¹ = OH, R¹² = H) with a lower alkyl Grignard or lithium reagent may yield compound I (R¹¹ = lower alkyl, R¹² = H).

SCHEME I

PREPARATION OF LACTONE INTERMEDIATES



- 25 -

SCHEME II

PREPARATION OF FINAL PRODUCTS

5

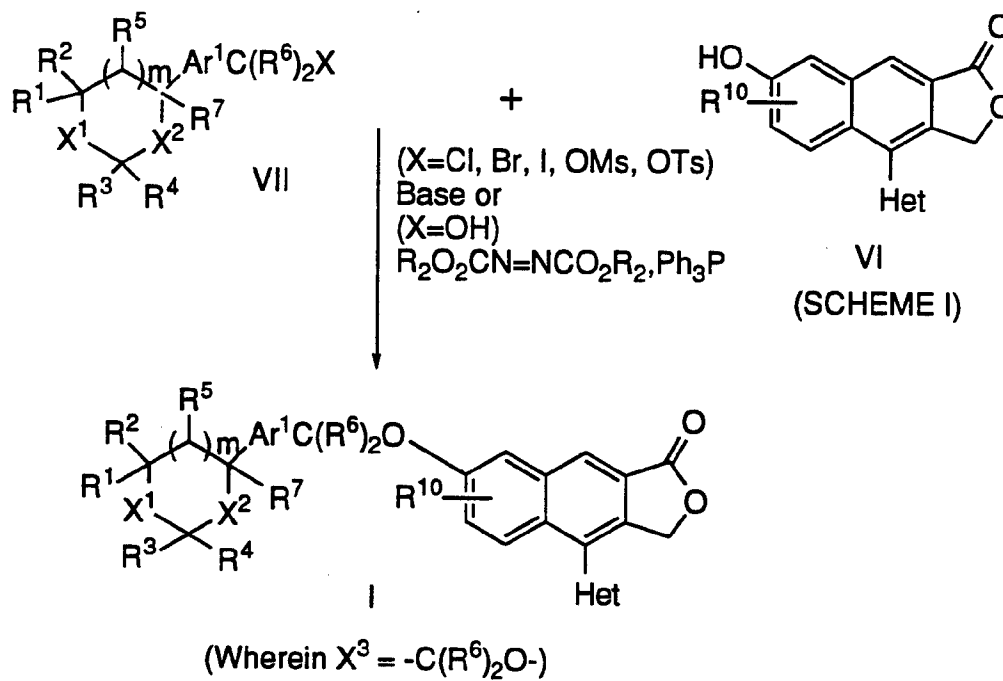
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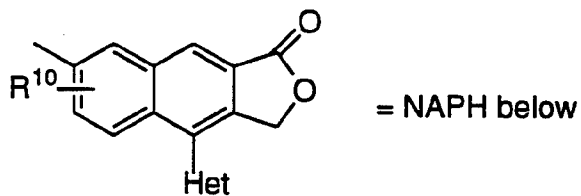


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SCHEME III

PREPARATION OF FINAL PRODUCTS

5



10

HO-NAPH
VI
(SCHEME I)



TfO-NAPH
VIII



MeO₂C-NAPH
IX

15



BrCH₂-NAPH
XII

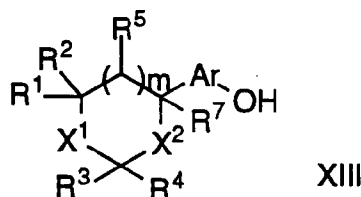


HOCH₂-NAPH
XI

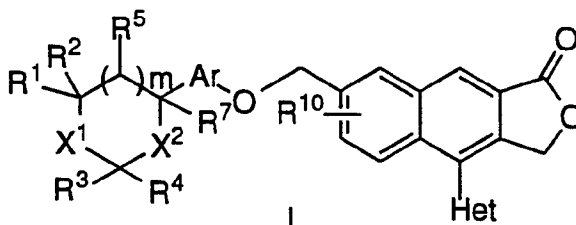
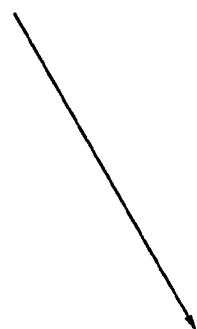


HO₂C-NAPH
X

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(Wherein X³ = -OCH₂-)

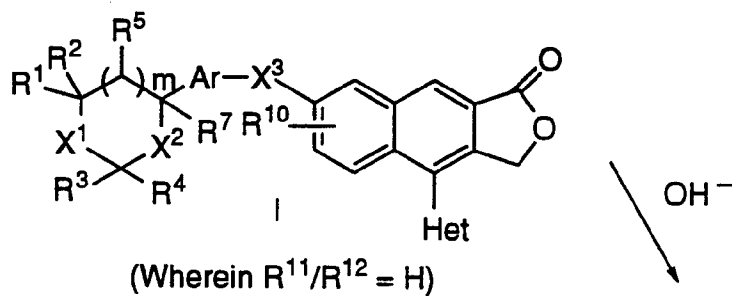
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SCHEME IV

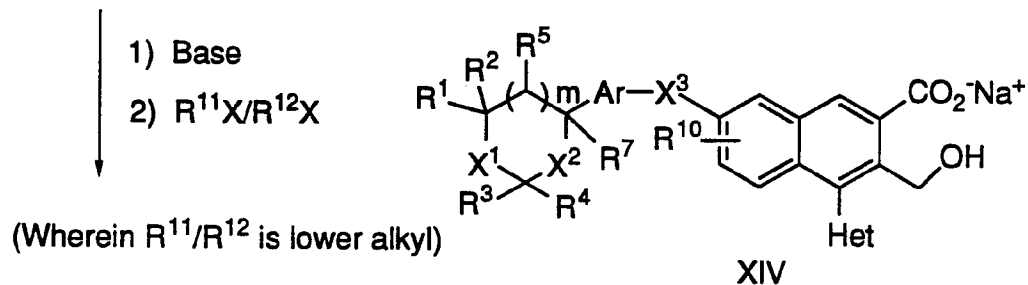
PREPARATION OF FINAL PRODUCTS

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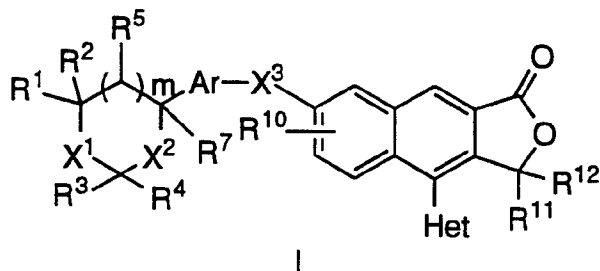
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PCC

(Wherein R^{11} is hydroxyl)
(and R^{12} is H)

25

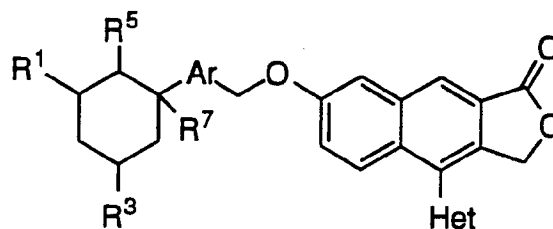
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Representative Compounds

Table I illustrates compounds representative of the present invention.

TABLE I

1b

EX.	R ¹	R ³	R ⁵	R ⁷	Ar	Het
1	H	H	H	OH	Phe	3-Fu
2	H	H	H	OMe	Phe	3-Fu
3	H	H	H	OH	Phe	3-Py
4	H	H	H	OH	Phe	3-Fu
5	-CH=CH-		H	OH	Phe	3-Fu
6	-CH ₂ CH ₂ -		H	OH	Phe	3-Fu
7	H	H	H	OH	5,3-Pye	3-Fu
8	-CH ₂ O-		H	OH	Phe	3-Fu
9	H	H	H	OH	Phe	3-Th
10	H	H	H	OH	5,3-Pye	3-Th
11	-CH ₂ O-		H	OH	Phe	3-Th
12	-CH ₂ O-		H	OMe	Phe	3-Th
13	-OCH ₂ -		H	OH	Phe	3-Fu
14	H	H	H	OH	6,2-Pye	3-Th
15	H	H	H	Et	Phe	3-Fu
16	-CH ₂ CH ₂ -		OMe	OH	Phe	3-Fu
17	H	H	H	OH	6,2-Pye	3-Fu

- 29 -

EX.	R ¹	R ³	R ⁵	R ⁷	Ar	Het
18	H	H	H	OH	2,4-Pye	3-Fu
19	H	H	H	OH	4,2-Pye	3-Fu
20	-CH ₂ O-		H	OH	6,2-Pye	3-Fu
21	-CH ₂ O-		H	OH	2,4-Pye	3-Fu
22	-CH ₂ O-		H	OH	4,2-Pye	3-Fu
23	-CH ₂ O-		H	OH	6,2-Pye	3-Fu

10

Assays for Determining Biological Activity

Compounds of Formula I can be tested using the following assays to determine their mammalian leukotriene biosynthesis inhibiting activity.

15

Human 5-Lipoxygenase Inhibitor Screen

Objective of the Assay: The objective of the assay is to select agents which specifically inhibit the activity of human 5-lipoxygenase using a 100,000x g supernatant fraction prepared from insect cells infected with recombinant baculovirus containing the coding sequence for human 5-lipoxygenase. Enzyme activity is measured spectrophotometrically from the optimal rate of conjugated diene formation (A₂₃₄) measured after the incubation of the enzyme with arachidonic acid in the presence of ATP, calcium ions, and phosphatidylcholine.

25

Description of Procedure: The activity of 5-lipoxygenase is measured using a spectrophotometric assay and recombinant human 5-lipoxygenase as a source of enzyme. The 100,000x g fraction from S19 cells infected with the recombinant baculovirus rvH5LO(8-1) containing the coding region sequence for human 5-lipoxygenase is prepared as described by Denis *et al.* (J. Biol. Chem., 266, 5072-79 (1991)). The enzymatic activity is measured, using a spectrophotometric assay from the optimal rate of conjugated diene formation (A₂₃₄) using the procedure described by Riendeau *et al.* (Biochem. Pharmacol., 38, 2323-2321, 1989) with minor modifications.

30

- 30 -

The incubation mixture contains 50 mM sodium phosphate pH 7.4, 0.2 mM ATP, 0.2 mM CaCl_2 , 20 μM arachidonic acid (5 μL from a 100-fold concentrated solution in ethanol), 12 $\mu\text{g/mL}$ phosphatidylcholine, an aliquot of the 100,000x g fraction (2-10 μL) and inhibitor (0.5 mL final volume). Inhibitors are added as 500-fold concentrated solutions in DMSO. Reactions are initiated by the addition of an aliquot of the enzyme preparation and the rate of conjugated diene formation is followed for 2 minutes at room temperature. The reactions are performed in semi-micro cuvettes (0.7 mL capacity, 10 mm path length, and 4 mm internal width) and the absorbance changes are recorded with a Hewlett-Packard diode array spectrophotometer (HP 8452A) connected to the ChemStation using UV/VIS Kinetics Software (Hewlett-Packard). Enzymatic activity is calculated from the optimal rate of the reaction by a linear fit of the variation of A_{234} during the first twenty seconds using the least square method for the equation $A_{234} = V_0 t + A_0$ where V_0 is the rate, t is the time and A_0 is the absorbance at zero time. The results are expressed as percentages of inhibition of the reaction rate relative to controls (typically between 0.15-0.21 AU/min) containing the DMSO vehicle.

Rat Peritoneal Polymorphonuclear (PMN) Leukocyte Assay

Rats under ether anesthesia are injected (i.p.) with 8 mL of a suspension of sodium caseinate (6 g in ca. 50 mL water). After 15-24 hr. the rats are sacrificed (CO_2) and the cells from the peritoneal cavity are recovered by lavage with 20 mL of buffer (Eagles MEM containing 30 mM HEPES adjusted to pH 7.4 with NaOH). The cells are pelleted (350x g, 5 min.), resuspended in buffer with vigorous shaking, filtered through lens paper, recentrifuged and finally suspended in buffer at a concentration of 10 cells/mL. A 500 μL aliquot of PMN suspension and test compound are pre-incubated for 2 minutes at 37° , followed by the addition of 10 μM calcium ionophore A-23187. The suspension is stirred for an additional 4 minutes then bioassayed for LTB_4 content by adding an aliquot to a second 500 μL portion of the PMN at 37°C . The LTB_4 produced in the first incubation causes aggregation of the second

- 31 -

PMN, which is measured as a change in light transmission. The size of the assay aliquot is chosen to give a submaximal transmission change (usually -70%) for the untreated control. The percentage inhibition of LTB₄ formation is calculated from the ratio of transmission change in the sample to the transmission change in the compound-free control.

Human Polymorphonuclear (PMN) Leukocyte LTB₄ Assay

A. Preparation of Human PMN. Human blood is obtained by antecubital venepuncture from consenting volunteers who have not taken medication within the previous 7 days. The blood is immediately added to 10% (v/v) trisodium citrate (0.13 M) or 5% (v/v) sodium heparin (1000 IU/mL). PMNs are isolated from anticoagulated blood by dextran sedimentation of erythrocytes followed by centrifugation through Ficoll-Hypaque (specific gravity 1.077), as described by Boyum, A., Scand. J. Clin. Lab. Invest., 21 (Supp 97), 77 (1968). Contaminating erythrocytes are removed by lysis following exposure to ammonium chloride (0.16 M) in Tris buffer (pH 7.65), and the PMNs resuspended at 5×10^5 cells/mL in HEPES (15 mM)-buffered Hanks balanced salt solution containing Ca²⁺ (1.4 mM) and Mg²⁺ (0.7 mM), pH 7.4. Viability is assessed by Trypan blue exclusion.

B. Generation and Radioimmunoassay of LTB₄. PMNs (0.5 mL; 2.5×10^5 cells) are placed in plastic tubes and incubated (37°C, 2 min) with test compounds at the desired concentration or vehicle (DMSO, final concentration 0.2%) as control. The synthesis of LTB₄ is initiated by the addition of calcium ionophore A23187 (final concentration 10 µM) or vehicle in control samples and allowed to proceed for 5 minutes at 37°C. The reactions are then terminated by the addition of cold methanol (0.25 mL) and samples of the entire PMN reaction mixture are removed for radioimmunoassay of LTB₄.

Samples (50 µL) of authentic LTB₄ of known concentration in radioimmunoassay buffer (RIA) buffer (potassium phosphate 1 mM; disodium EDTA 0.1 mM; Thimerosal 0.025 mM; gelatin 0.1%, pH 7.3) or PMN reaction mixture diluted 1:1 with RIA buffer are added to reaction tubes. Thereafter [³H]-LTB₄ (10 nCi in

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100 μ L RIA buffer) and LTB₄-antiserum (100 μ L of a 1:3000 dilution in RIA buffer) are added and the tubes vortexed. Reactants are allowed to equilibrate by incubation overnight at 4°C. To separate antibody-bound from free LTB₄, aliquots (50 μ L) of activated charcoal (3% activated charcoal in RIA buffer containing 0.25% Dextran T-70) are added, the tubes vortexed, and allowed to stand at room temperature for 10 minutes prior to centrifugation (1500x g; 10 min; 4°C). The supernatants containing antibody-bound LTB₄ are decanted into vials and Aquasol 2 (4 mL) is added. Radioactivity is quantified by liquid scintillation spectrometry. The specificity of the antiserum and the sensitivity of the procedure have been described by Rokach *et al.* Prostaglandins Leukotrienes and Medicine, 13, 21 (1984). The amount of LTB₄ produced in test and control samples is calculated. Inhibitory dose-response curves are constructed using a four-parameter algorithm and from these the IC₅₀ values are determined.

Asthmatic Rat Assay

Rats are obtained from an inbred line of asthmatic rats. Both female (190-250 g) and male (260-400 g) rats are used. Egg albumin (EA), grade V, crystallized and lyophilized, is obtained from Sigma Chemical Co., St. Louis. Aluminum hydroxide is obtained from the Regis Chemical Company, Chicago. Methysergide bimalate is supplied by Sandoz Ltd., Basel.

The challenge and subsequent respiratory recordings are carried out in a clear plastic box with internal dimensions 10x6x4 inches. The top of the box is removable; in use, it is held firmly in place by four clamps and an airtight seal is maintained by a soft rubber gasket. Through the center of each end of the chamber a DeVilbiss nebulizer (No. 40) is inserted via an airtight seal and each end of the box also has an outlet. A Fleisch No. 0000 pneumotachograph is inserted into one end of the box and coupled to a Grass volumetric pressure transducer (PT5-A) which is then connected to a Beckman Type R Dynograph through appropriate couplers. While aerosolizing the antigen, the outlets are open and the pneumotachograph is isolated

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from the chamber. The outlets are closed and the pneumotachograph and the chamber are connected during the recording of the respiratory patterns. For challenge, 2 mL of a 3% solution of antigen in saline is placed into each nebulizer and the aerosol is generated with air from a small Potter diaphragm pump operating at 10 psi and a flow of 8 L/minute.

Rats are sensitized by injecting (subcutaneously) 1 mL of a suspension containing 1 mg EA and 200 mg aluminum hydroxide in saline. They are used between days 12 and 24 postsensitization. In order to eliminate the serotonin component of the response, rats are pretreated intravenously 5 minutes prior to aerosol challenge with 3.0 mg/kg of methysergide. Rats are then exposed to an aerosol of 3% EA in saline for exactly 1 minute, then their respiratory profiles are recorded for a further 30 minutes. The duration of continuous dyspnea is measured from the respiratory recordings.

Compounds are generally administered either orally 1-4 hours prior to challenge or intravenously 2 minutes prior to challenge. They are either dissolved in saline or 1% methocel or suspended in 1% methocel. The volume injected is 1 mL/kg (intravenously) or 10 mL/kg (orally). Prior to oral treatment rats are starved overnight. Their activity is determined in terms of their ability to decrease the duration of symptoms of dyspnea in comparison with a group of vehicle-treated controls. Usually, a compound is evaluated at a series of doses and an ED50 is determined. This is defined as the dose (mg/kg) which would inhibit the duration of symptoms by 50%.

Pulmonary Mechanics in Trained Conscious Squirrel Monkeys

The test procedure involves placing trained squirrel monkeys in chairs in aerosol exposure chambers. For control purposes, pulmonary mechanics measurements of respiratory parameters are recorded for a period of about 30 minutes to establish each monkey's normal control values for that day. For oral administration, compounds are dissolved or suspended in a 1% methocel solution (methylcellulose, 65HG, 400 cps) and given in a volume of 1 mL/kg body weight. For

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aerosol administration of compounds, a DeVilbiss ultrasonic nebulizer is utilized. Pretreatment periods vary from 5 minutes to 4 hours before the monkeys are challenged with aerosol doses of Ascaris antigen.

5 Following challenge, each minute of data is calculated by computer as a percent change from control values for each respiratory parameter including airway resistance (RL) and dynamic compliance (Cdyn). The results for each test compound are subsequently obtained for a minimum period of 60 minutes post challenge which are then compared to previously obtained historical baseline control values for
10 that monkey. In addition, the overall values for 60 minutes post-challenge for each monkey (historical baseline values and test values) are averaged separately and are used to calculate the overall percent inhibition of Ascaris antigen response by the test compound. For
15 statistical analysis, paired t-test is used. (See: McFarlane, C.S. et al., Prostaglandins, 28:173-182, 1984, and McFarlane, C.S. et al., Agents Actions 22:63-68, 1987.)

Prevention of Induced Bronchoconstriction in Allergic Sheep

20 A. Rationale. Certain allergic sheep with known sensitivity to a specific antigen (Ascaris suum) respond to inhalation challenge with acute and late bronchial responses. The time course of both the acute and the late bronchial responses approximates the time course observed in asthmatics and the pharmacological modification of both responses is similar to that found in man. The effects of antigen in
25 these sheep are largely observed in the large airways and are conveniently monitored as changes in lung resistance and specific lung resistance.

30 B. Methods. Animal Preparation: Adult sheep with a mean weight of 35 kg (range, 18 to 50 kg) are used. All animals used meet two criteria: a) they have a natural cutaneous reaction to 1:1,000 or 1:10,000 dilutions of Ascaris suum extract (Greer Diagnostics, Lenois, NC) and b) they have previously responded to inhalation challenge with Ascaris suum with both an acute bronchoconstriction and

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a late bronchial obstruction (Abraham, W.M. *et al.*, Am. Rev. Resp. Dis., 128, 839-44 (1983)).

5 Measurement of Airway Mechanics: The unsedated sheep
are restrained in a cart in the prone position with their heads
immobilized. After topical anesthesia of the nasal passages with 2%
lidocaine solution, a balloon catheter is advanced through one nostril
into the lower esophagus. The animals are then intubated with a cuffed
10 endotracheal tube through the other nostril using a flexible fiberoptic
bronchoscope as a guide. Pleural pressure is estimated with the
esophageal balloon catheter (filled with one mL of air), which is
positioned such that inspiration produces a negative pressure deflection
with clearly discernible cardiogenic oscillations. Lateral pressure in the
trachea is measured with a sidehole catheter (inner dimension, 2.5 mm)
15 advanced through and positioned distal to the tip of the nasotracheal
tube. Transpulmonary pressure, the difference between tracheal
pressure and pleural pressure, is measured with a differential pressure
transducer (DP45; Validyne Corp., Northridge, CA). For the
measurement of pulmonary resistance (R_L), the maximal end of the
20 nasotracheal tube is connected to a pneumotachograph (Fleisch, Dyna
Sciences, Blue Bell, PA). The signals of flow and transpulmonary
pressure are recorded on an oscilloscope (Model DR-12; Electronics for
Medicine, White Plains, NY) which is linked to a PDP-11 Digital
computer (Digital Equipment Corp., Maynard, MA) for on-line
25 calculation of R_L from transpulmonary pressure, respiratory volume
obtained by integration and flow. Analysis of 10-15 breaths is used for
the determination of R_L . Thoracic gas volume (V_{tg}) is measured in a
body plethysmograph, to obtain specific pulmonary resistance ($SRL =$
30 $R_L \cdot V_{tg}$).

Aerosol Delivery Systems: Aerosols of Ascaris suum
extract (1:20) are generated using a disposable medical nebulizer
(Raindrop®, Puritan Bennett), which produces an aerosol with a mass
median aerodynamic diameter of 6.2 μ M (geometric standard deviation,
2.1) as determined by an electric size analyzer (Model 3030; Thermal

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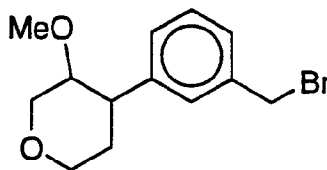
Systems, St. Paul, MN). The output from the nebulizer is directed into a plastic t-piece, one end of which is attached to the nasotracheal tube, the other end of which is connected to the inspiratory part of a Harvard respirator. The aerosol is delivered at a tidal volume of 500 mL of a rate of 20 per minute. Thus, each sheep receives an equivalent dose of antigen in both placebo and drug trials.

Experimental Protocol: Prior to antigen challenge baseline measurements of SRL are obtained, infusion of the test compound is started 1 hr prior to challenge, the measurement of SRL repeated and then the sheep undergoes inhalation challenge with Ascaris suum antigen. Measurements of SRL are obtained immediately after antigen challenge and at 1, 2, 3, 4, 5, 6, 6.5, 7, 7.5, and 8 hrs after antigen challenge. Placebo and drug tests are separated by at least 14 days. In a further study, sheep are given a bolus dose of the test compound followed by an infusion of the test compound for 0.5-1 hr prior to Ascaris challenge and for 8 hrs after Ascaris as described above.

Statistical Analysis: A Kruskal-Wallis one way ANOVA test is used to compare the acute immediate responses to antigen and the peak late response in the controls and the drug treated animals.

PREPARATION OF BENZYL HALIDES

Halide 1: 3-[4-(4-Methoxy)tetrahydropyranyl]benzyl bromide



Step 1: 3-[4-(4-Hydroxy)tetrahydropyranyl]toluene

To a solution of 3-bromotoluene (24.3 mL; Aldrich) in THF (250 mL) stirred at -78°C was added a solution of n-BuLi in hexane (1.75 M; 114 mL). After 45 min., the resulting white suspension was treated with a solution of tetrahydropyran-4-one (18.5 mL; Aldrich) in THF (125 mL). After 45 min. at -78°C, the mixture

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was stirred for 1.5 hr. at room temperature (r.t.) Saturated aqueous NH₄Cl was then added and the organic phase separated. The aqueous phase was extracted with EtOAc. The combined organic phases were washed with brine, dried (MgSO₄) and evaporated. Flash chromatography of the residue (silica gel; hexane/EtOAc (1:1)) followed by crystallization in hexane/EtOAc afforded the title compound as a white solid.

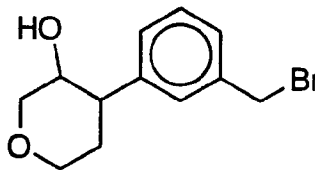
Step 2: 3-[4-(4-Methoxy)tetrahydropyranyl]toluene

To a 0°C solution of the alcohol from Step 1 (38 g) in DMF (300 mL) were added NaH (60% in mineral oil; 16 g) and methyl iodide (31 mL). The mixture was stirred under nitrogen at r.t. for 15 hr. before H₂O (1 L) was added. The aqueous phase was extracted with EtOAc and the combined organic phases were washed with brine, dried (MgSO₄) and evaporated. Flash chromatography of the residue (silica gel; hexane/EtOAc (4:1)) yielded the title ether as a colorless liquid.

Step 3: 3-[4-(4-Methoxy)tetrahydropyranyl]benzyl bromide

A mixture of the toluene (16 g) from Step 2, N-bromosuccinimide (14.6 g) and azoisobutyronitrile (AIBN) (127 mg) in CC1₄ (250 mL) was refluxed for 1.5 hr. Filtration and evaporation of the filtrate gave the desired benzyl bromide.

Halide 2: 3-[4-(4-Hydroxy)tetrahydropyranyl]benzyl bromide

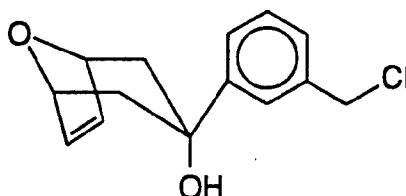


Following the procedure described for Halide 1, Step 3, but substituting 3-[4-(4-hydroxy)tetrahydropyranyl]toluene (from Halide 1, Step 1) for 3-[4-(4-methoxy)tetrahydropyranyl]toluene, the title product was obtained as a yellow solid.

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Halide 3: 3-[3-(3 α -Hydroxy-8-oxabicyclo[3.2.1]oct-6-enyl)]benzyl chloride

5



Step 1: 3-Bromo-O-tetrahydropyranylbenzyl alcohol

10

To a solution of 3-bromobenzyl alcohol (11.5 g; Aldrich) dissolved in CH₂Cl₂ (100 mL) at 0°C and p-toluenesulfonic acid monohydrate (116 mg) was added 6.2 mL of DHP. The resulting solution was stirred at r.t. for 3 hr. then was quenched with NH₄OAc. The aqueous phase was extracted with CH₂Cl₂. The combined organic phases were washed with brine, dried (MgSO₄) and evaporated. Flash chromatography of the residue (silica gel: hexane/EtOAc (9:1)) afforded the title compound as an oil.

15

Step 2: 3-[3-(3 α -Hydroxy-8-oxabicyclo[3.2.1]oct-6-enyl)-O-tetrahydropyranyl]benzyl alcohol

20

Following the procedure described in Halide 1, Step 1, but substituting 3-bromo-O-tetrahydro-pyranylbenzyl alcohol from Step 1 for 3-bromotoluene, and substituting 8-oxabicyclo[3.2.1]oct-6-en-3-one (J. Amer. Chem. Soc., 100, 1765 (1978)) for tetrahydropyran-4-one, the title compound was obtained.

25

Step 3: 3-[3-(3 α -Hydroxy-8-oxabicyclo[3.2.1]oct-6-enyl)]benzyl alcohol

30

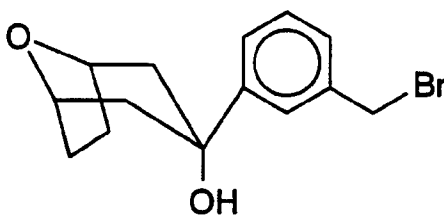
A solution of the THP derivative (600 mg) from Step 2, in 8 mL of EtOH and pyridinium p-toluenesulfonate (18 mg) was heated at 55°C for 3 hours. Evaporation of the solvent followed by a flash column (hexane/EtOAc, 1:1) gave the title compound.

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Step 4: 3-[3-(3 α -Hydroxy-8-oxabicyclo[3.2.1]oct-6-enyl)benzyl chloride

To a solution of the alcohol (300 mg, 1.3 mmol) obtained from Step 3, in CH₂Cl₂ (5 mL) and CCl₄ (144 μ L) was added Ph₃P (407 mg). The reaction was heated overnight at 70°C. The reaction was diluted with 10% EtOAc in hexane and flash chromatographed directly to give the title compound.

Halide 4: 3-[3-(3 α -Hydroxy-8-oxabicyclo[3.2.1]-octanyl)]benzyl bromide



Step 1: 3-[3-(3 α -Hydroxy-8-oxabicyclo[3.2.1]oct-6-enyl)toluene

Following the procedure described in Halide 1, Step 1, but substituting 8-oxabicyclo[3.2.1]oct-6-en-3-one (J.A.C.S., 100, 1765 (1978)) for tetrahydropyran-4-one, the title compound was obtained.

Step 2: 3-[3-(3 α -Hydroxy-8-oxabicyclo[3.2.1]octan-yl)toluene

A mixture of the bicyclic alkene (1.50 g) from Step 1 and 10% Pd/C (0.5 g) in EtOAc (150 mL) and MeOH (15 mL) was stirred under hydrogen at r.t. for 18 hr. The reaction mixture was then filtered over celite and evaporation of the filtrate afforded a yellow waxy solid. Trituration in ether followed by filtration and evaporation of the filtrate then gave a white solid. Purification by flash chromatography (silica gel; hexane/EtOAc, 3:1) yielded the title toluene.

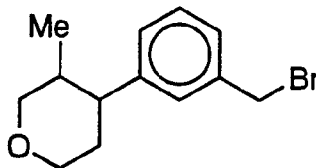
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Step 3: 3-[3-(3 α -Hydroxy-8-oxabicyclo[3.2.1]octan-yl)benzyl bromide

5 A mixture of the toluene (0.39 g) from Step 2, NBS (0.33 g) and benzoyl peroxide (10 mg) in CCl₄ (10 mL) was refluxed and irradiated with visible light for 1.5 hr. Filtration and evaporation of the filtrate gave the title benzyl bromide.

Halide 5: 3-[4-(4-Methyl)tetrahydropyranyl]benzyl bromide

10



15 Step 1: 3-[4-(4-Chloro)tetrahydropyranyl]toluene

A solution of 3-[4-(4-hydroxy)tetrahydro-pyranyl]toluene (Halide 1, Step 1) 0.57 g in CHCl₃ (10 mL) was added to a slurry of P₂Cl₅ (1.92 g) and K₂CO₃ (0.41 g) in CHCl₃ (10 mL) stirred at 0°C for 35 min. The reaction mixture was filtered and evaporation of the
20 filtrate afforded the title compound as a yellow semi-solid.

Step 2: 3-[4-(4-Methyl)tetrahydropyranyl]toluene

To a 0°C solution of the chloride (452 mg, Step 1) in CH₂Cl₂ (7 mL) was added a solution of trimethylaluminum in hexane (2.0M; 2.2 mL; Aldrich). The mixture was stirred at r.t. for 19 hr.
25 before saturated NaHCO₃ was added. The aqueous phase was extracted twice with CH₂Cl₂ and the combined organic phases were dried (MgSO₄) and evaporated. Flash chromatography of the residue (silica gel; hexane/EtOAc, 9:1) yielded the title compound as a colorless liquid.

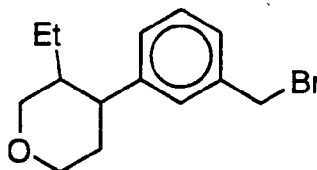
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Step 3: 3-[4-(4-Methyl)tetrahydropyranyl]benzyl bromide

Following the procedure described in Halide 1, Step 3, but substituting the toluene from Step 2 for 3-[4-(4-methoxy)tetrahydro-pyranyl]toluene, the title product was obtained as an oil.

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Halide 6: 3-[4-(4-Ethyl)tetrahydropyranyl]benzyl bromide

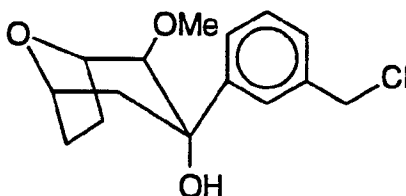


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Following the procedure described in Halide 5, Steps 2-3, but substituting triethylaluminum for trimethylaluminum, the title benzyl bromide was obtained as an oil.

10

Halide 7: 3-[3-(3 α -Hydroxy-2-methoxy-8-oxabicyclo-[3.2.1]-octanyl)]benzyl chloride



15

Step 1: 3-[3-(3 α -Hydroxy-2-methoxy-8-oxabicyclo[3.-2.1]octan-6-enyl)-O-tetrahydropyranyl]benzyl alcohol

20

Following the procedure described in Halide 1, Step 1, but substituting 3-bromo-O-tetrahydro-pyranyl benzyl alcohol (from Halide 3, Step 1) for 3-bromotoluene and substituting 2-methoxy-8-oxabicyclo[3.2.1]oct-6-en-3-one (Tet. Lett., 1990, 31, 4109) for tetrahydropyran-4-one, the title compound was obtained.

25

Step 2: 3-[3-(3 α -Hydroxy-2-methoxy-8-oxabicyclo[3.-2.1]octanyl)-O-tetrahydropyranyl]benzyl alcohol

30

Following the procedure described in Halide 4, Step 2, but substituting the bicyclic alkene derivative from Step 1 for 3-[3-(3 α -hydroxy-8-oxabicyclo[3.2.1]oct-6-enyl)-O-tetrahydropyranyl]benzyl alcohol, the title compound was obtained.

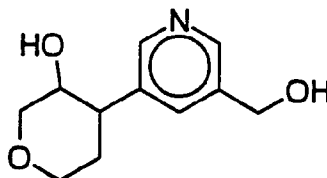
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Step 3: 3-[3-(3 α -Hydroxy-2-methoxy-8-oxabicyclo[3.2.1]octanyl)]-benzyl chloride

Following the procedure described in Halide 3, Steps 3-4, but substituting the THP derivative from Step 2 for 3-[3-(3 α -hydroxy-8-oxabicyclo-[3.2.1]octanyl)-O-tetrahydropyranyl]benzyl alcohol, the title compound was obtained.

PREPARATION OF ALCOHOLS

Alcohol 1: 5-[4-(4-Hydroxy)tetrahydropyranyl]-pyridin-3-ylmethanol



Step 1: 5-Bromo-0-tert-butyl-diphenylsilylpyridin-3-ylmethanol

To a solution of 5-bromopyridin-3-ylmethanol (Chem. Pharm. Bull. 1990, 38, 2446) (29 g, 154 mmol) and tert-butylchlorodiphenylsilane (47.5 g, 173 mmol) in CH₂Cl₂ (500 mL) at r.t., there was added imidazole (15.8 g, 232 mmol). The mixture was stirred for 1 hr. and filtered. The filtrate was evaporated and the residue chromatographed on silica gel eluting with a 1:7 mixture of EtOAc and hexane, to afford the product as a colorless oil.

Step 2: 5-[4-(4-Hydroxy)tetrahydropyranyl]-0-tert-butyl-diphenylsilylpyridin-3-ylmethanol

To a solution of the silylether from Step 1 (50 g, 117 mmol) in THF (500 mL), cooled to -70°C, there was slowly added n-BuLi 1.12 M in hexanes (115 mL, 129 mmol) affording a dark brown solution. To this, there was added a solution of tetrahydro-4H-pyran-4-one (14.1 g, 141 mmol) in THF (925 mL). The resulting mixture was stirred for 1 hr. at -70°C, then quenched slowly with saturated aqueous NH₄Cl (50 mL) and allowed to warm up to r.t. After diluting with EtOAc (500 mL) the mixture was washed (4x) with brine,

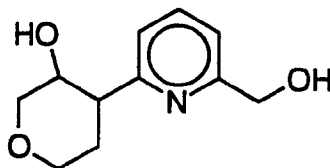
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dried over Na₂SO₄ and evaporated. Chromatography on silica gel, eluting with EtOAc, afforded the product as an oil which solidified.

5 **Step 3: 5-[4-(4-Hydroxy)tetrahydropyranyl]pyridin-3-ylmethanol**

To a solution of the silylether from Step 2 (20.35 g, 45.5 mmol) in THF (350 mL), there was added Bu₄NF 1M in THF (52 mL) and the mixture was stirred at r.t. for 1 hr. The solvent was evaporated and the residue chromatographed as a short column of silica gel, eluting with a 1:4 mixture of EtOH and EtOAc to afford the title product which was obtained, after trituration with Et₂O and filtration, as a light yellow solid; m.p. 145-147°C.

15 **Alcohol 2: 6-[4-(4-Hydroxy)tetrahydropyranyl]-pyridin-2-ylmethanol**



20 **Step 1: 2-Bromo-6-[4-(4-hydroxy)tetrahydropyranyl]-pyridine**

A solution of 2,6-dibromopyridine (15 g) in Et₂O (375 mL) was cooled to -78°C. To the resulting suspension was slowly added n-BuLi 2M in hexanes (47.5 mL, 0.9 eq.) and the resulting mixture was stirred for a further 15 min. at -78°C. There was slowly added a solution of tetrahydro-4H-pyran-4-one (11.6 g) in Et₂O (25 mL). The resulting white suspension was stirred at -78°C for an additional 15 min. There was added saturated aqueous NH₄Cl (100 mL) and the mixture was allowed to warm up to r.t. After dilution with EtOAc, the organic phase was washed (4x) with brine, dried and evaporated. The residue was trituated with Et₂O and filtered to afford the title product as a white solid; m.p. 131-133°C.

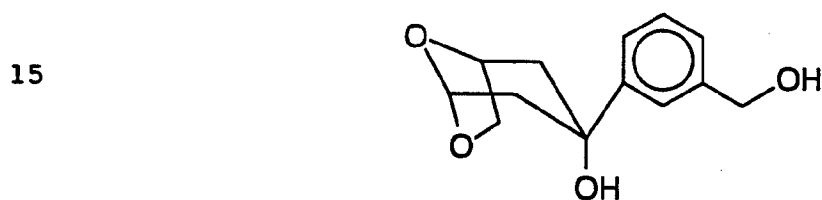
30 **Step 2: 6-[4-(4-Hydroxy)tetrahydropyranyl]pyridin-2-ylmethanol**

To a solution of the bromo derivative from Step 1 (7.7 g) in THF (50 mL) and Et₂O (150 mL), cooled to 0°C, there was slowly

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added n-BuLi 2M in hexanes (30 mL) affording a red-brown suspension. An inlet tube above the surface of the mixture was connected to a flask in which paraformaldehyde (25 g) was gently heated at 175°C to generate formaldehyde. When all the
5 paraformaldehyde had been decomposed, to the reaction mixture was added saturated aqueous NH₄Cl (100 mL) and EtOAc (500 mL). The organic phase was washed (4x) with brine, dried and evaporated to a residue which was chromatographed on silica gel, eluting with EtOAc to
10 afford the title product as a thick yellow oil.

Alcohol 3: (1S,5R)3-[3-(3 α -Hydroxy-6,8-dioxabicyclo[3.2.1]-octanyl)]benzyl alcohol



Step 1: 3-Bromo-O-tert-butyldiphenylsilylbenzyl alcohol

20 To a solution of 3-bromobenzyl alcohol (25 g, 134 mmol) in anhydrous DMF (300 mL) was added triethylamine (17.6 g, 174 mmol) followed by t-butyldiphenylsilyl chloride (40.4 g, 147 mmol). The mixture was stirred for 24 hr., poured into a saturated aqueous NH₄Cl solution (1 L), and extracted with Et₂O. The combined organic
25 layers were washed with brine, dried over MgSO₄ and evaporated. Flash chromatography on silica gel (2.5% EtOAc in hexane) afforded the title compound as a colorless oil.

Step 2: 2,4-Di-O-p-toluenesulfonyl-1,6-anhydro- β -D-glucose

30 To a solution of 1,6-anhydro- β -D-glucose (50 g, 308 mmol) in dry pyridine (100 mL) at 0°C was added dropwise to a solution of p-toluenesulfonyl chloride (123 g, 647 mmol) dissolved in CHCl₃ (350 mL) and pyridine (200 mL). The reaction mixture was stirred at r.t. for at least 2 days. Water was added and the reaction mixture was stirred for ~1 hr, then the organic layer was decanted and

- 45 -

the aqueous phase was re-extracted with CHCl_3 . The combined organic layers were washed with H_2SO_4 (10%) until the pH remained acidic, then finally washed with a saturated NH_4OAc solution. The resulting organic layer was dried over MgSO_4 and the solvent evaporated. The syrup obtained was flash chromatographed on silica gel eluting with hexane:EtOAc (1:1) to give the title compound as an oil.

Step 3: (1S,3S,5R) 6,8-Dioxabicyclo[3.2.1]octan-3-ol

The ditosylate derivative from Step 1 (107 g, 0.228 mmol) was dissolved in THF (1.6 L) at -40°C and Super-Hydride in THF (800 mL, 1 M, 0.8 mmol) was slowly added. The resulting reaction mixture was stirred at r.t. overnight. The reaction was cannulated into cold H_2O (226 mL) using external cooling, then NaOH 3N (640 mL, 1.92 mmol) and H_2O_2 (30%) (490 mL, 4.3 mmol) were successively added. The reaction was stirred at r.t. for 1 hr. Then the supernatant (THF layer) was separated from the aqueous layer and concentrated. The resulting residue was combined with the aqueous layer and extracted with CH_2Cl_2 using a continuous extractor. The organic layer was dried (MgSO_4) and evaporated to dryness. The oily residue was dissolved in hot Et_2O , filtered and evaporated to dryness affording the title compound contaminated with the 2-octanol isomer. The crude product was used as such for the next step.

Step 4: (1S,5R) 6,8-Dioxabicyclo[3.2.1]octan-3-one

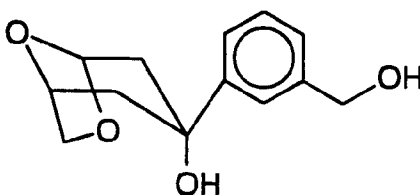
The crude alcohol from Step 2 (16.6 g, 89 mmol) in CH_2Cl_2 (200 mL) was added slowly to a suspension of PCC (38.4 g, 178 mmol) and celite (22 g) in CH_2Cl_2 (400 mL) and stirred for 1 hr. The reaction mixture was diluted with Et_2O (600 mL) and filtered over celite. The filtrate was evaporated and the residue distilled with a Kugelrohr apparatus (100°C , 1.8 mm/Hg) affording the title product as an oil.

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Step 5: (1S,5R) 3-[3-(3 α -Hydroxy-6,8-dioxabicyclo-[3,2,1]-octanyl)benzyl alcohol

Following the procedure described in Halide 1, Step 1, but substituting 3-bromo-O-tert-butyl-diphenylsilylbenzyl alcohol (from Step 1) for 3-bromotoluene, and (1S,5R) 6,8-dioxabicyclo[3.2.1]-octan-3-one (from Step 4) for tetrahydropyran-4-one, the tert-butyl-diphenylsilylether derivative of the title compound was obtained. The crude product was treated with 1 equivalent of Bu₄NF in dry THF at r.t. for 1.5 hr. After evaporation of the solvent, the crude product was flash chromatographed on silica gel (hexane:EtOAc, 4:1) to afford the pure title product as a colorless oil.

Alcohol 4: (1R,5S) 3-[3-(3 α -Hydroxy-6,8-dioxabicyclo[3.2.1]-octanyl)benzyl alcohol



Step 1: (R)3-tert-Butyldimethylsilyloxy-1,2-epoxy-propane

To a 0°C mixture of tert-butyldimethylsilyl chloride (12.2 g) and imidazole (11.5 g) in CH₂Cl₂ (100 mL) was slowly added (S)-Glycidol (5.0 g; Aldrich). After 2.5 h. at 0°C, the reaction mixture was treated with saturated aqueous NH₄Cl and the organic phase was separated. The aqueous phase was extracted with CH₂Cl₂ (100 mL) and the combined organic phases were washed with H₂O and brine and then dried (MgSO₄) and evaporated. Flash chromatography of the residue (silica gel; hexane/EtOAc, 9:1) afforded the title epoxide as a colorless liquid.

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Step 2: (2R) 6,6-Dimethoxy-4-(1,3-propanediylthio)-1-tert-butyl
dimethylsilyloxyhexan-2-ol

To a solution of 2-(2,2-dimethoxyethyl)-1,3-dithiane (10.4 g; Liebigs Ann. Chem. 1989, 1045) in THF (50 mL) stirred at -30°C was added a solution of n-BuLi in hexane (1.3M; 38.5 mL; Aldrich). After 1 hr., the resulting purple solution was treated with a solution of the epoxide (9.4 g) from Step 1 in THF (10 mL). After 3 hr at -30°C, the reaction mixture was quenched with saturated aqueous NH₄Cl. Water was then added and the aqueous phase was extracted with Et₂O (2x). The combined organic phases were washed with H₂O and brine, dried (MgSO₄) and evaporated. Flash chromatography of the residue (silica gel; hexane/EtOAc, 4:1) afforded the title alcohol as a yellow liquid.

Step 3: (2R) 6,6-Dimethoxy-4-(1,3-propanediylthio)hexan-1,2-diol

To a solution of the silyl ether from Step 2 (9.94 g) in THF (50 mL) was added a solution of Bu₄NF in THF (1.0M; 26.5 mL; Aldrich). After 90 min. at r.t., the volatiles were evaporated. Flash chromatography of the residue (silica gel; hexane/EtOAc, 1:3) afforded the title diol as an amber oil.

Step 4: (2R) 6-Hydroxy-2-hydroxymethyl-3-(1,3-propanediylthio)tetrahydropyran

A solution of the diol from Step 3 (3.11 g) in THF (40 mL) and aqueous HCl (5% v/v; 25 mL) was stirred at 60°C for 18 hr. Water was then added and the aqueous phase was extracted with EtOAc (3x). The combined organic phases were washed with brine, dried (Na₂SO₄) and evaporated to afford the crude title compound as an amber gum.

Step 5: (2R) 6-Hydroxy-3-(1,3-propanediylthio)-p-toluenesulfonylmethyl tetrahydropyran

p-Toluenesulfonyl chloride (1.98 g; Aldrich) was added to a 0°C solution of the alcohol from Step 4 (2.05 g) in pyridine (10 mL).

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The mixture was stirred at r.t. for 90 min. before MeOH (1 mL) and CHCl₃ (50 mL) were added. The organic phase was washed with H₂O (2x), dried (MgSO₄) and evaporated. Flash chromatography of the residue (silica gel; hexane/EtOAc, 3:2) yielded the title tosylate as an
5 amber semi-solid.

Step 6: (1R,5S) 3-(1,3-Propanediylthio)-6,8-dioxabicyclo-
[3.2.1]octane

1,8-Diazabicyclo[5.4.0]undec-7-ene (1.5 mL; Aldrich) was
10 added to a solution of the tosylate from Step 5 (1.95 g) in CH₂Cl₂ (30 mL). The mixture was stirred at r.t. for 16 hr. before the volatiles were evaporated. Flash chromatography of the residue (silica gel; hexane/EtOAc, 3:1) gave the title compound as a white solid.

15 Step 7: (1R,5S) 6,8-Dioxabicyclo[3.2.1]octan-6-one

To a vigorously stirred mixture of the dithioketal from Step 6 (0.76 g) and CaCO₃ (0.42 g) in THF (12 mL) and H₂O (2.3 mL) was added in 10 min. a solution of Mercury (II) perchlorate trihydrate (1.73 g; Aldrich) in H₂O (1 mL). The mixture was stirred for 40 min.
20 at r.t. before another portion of mercury salt solution (0.56 g in 0.2 mL of H₂O) was slowly added. After 15 min., Et₂O (50 mL) was added and the reaction mixture was filtered. The organic filtrate was dried (MgSO₄) and evaporated to afford the title ketone as an amber liquid.

25 Step 8: (1R,5S) 3-[3-(3 α -Hydroxy-6,8-dioxabicyclo-
[3.2.1]octanyl)]benzyl alcohol

Following the procedure described in Alcohol 3, Step 5, but substituting 3-bromo-O-tert-butyl diphenylsilyl benzyl alcohol (from Alcohol 3, Step 1) for 3-bromotoluene and (1R,5S) 6,8-dioxabicyclo-
30 [3.2.1]octane-3-one (from Step 7) for tetrahydropyran-4-one, the tert butyldiphenylsilylether derivative of the title compound was obtained. The crude product was treated with 1 equivalent of Bu₄NF in dry THF at r.t. for 1.5 hr. After evaporation of the solvent, the crude product

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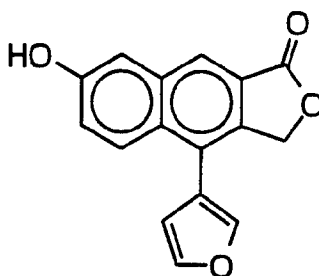
was flash chromatographed on silica gel (hexane:EtOAc, 4:1) to afford the pure title product as a colorless oil.

PREPARATION OF LACTONES

5

Lactone 1: 3-Hydroxymethyl-4-(3-furyl)-7-hydroxy-2-naphthoic acid, lactone form

10



15

Step 1: 3-Furaldehyde phenyl dithioacetal

To a solution of 3-furaldehyde (28.8 g) and thiophenol (69.2 g) in isopropyl acetate (300 mL), cooled in an ice-water bath, there was slowly added $\text{BF}_3 \cdot \text{Et}_2\text{O}$ (42.6 g). The resulting mixture was stirred in the cold for an additional hour. There was slowly added 10% aqueous K_2CO_3 (200 mL) and the phases were separated. The aqueous phase was extracted with EtOAc and the combined organic layers were washed with 10% aq. K_2CO_3 , then with H_2O (3x), dried and evaporated to an oil, which was used as such.

25

Step 2: 3-[3-Furyl-bis(phenylthio)methyl]2-(3-benzyloxy- α -hydroxybenzyl)butyrolactone

To a solution of the dithioacetal from Step 1 (60.0 g) in THF (500 mL), at -70°C there was slowly added 2.1M n-BuLi in hexanes (100 mL). The resulting dark solution was stirred for a further 20 min. at -70°C , then there was added dropwise 2-(5H) furanone (Omega Inc., 20.16 g), and 30 min. later, a solution of 3-benzyloxybenzaldehyde (Aldrich, 43.5 g) in THF (175 mL). The mixture was stirred a further hour at -70°C , then there was slowly added glacial HOAc (27 g). The mixture was allowed to warm up to

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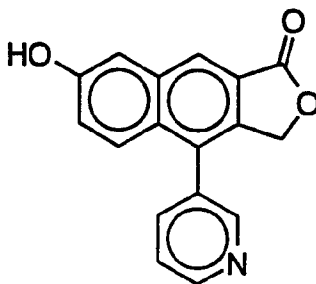
r.t., diluted with Et₂O (500 mL), washed with brine (4x), dried and evaporated. The residue was used as such in the next step.

5 Step 3: 3-Hydroxymethyl-4-(3-furyl)-7-hydroxy-2-naphthoic acid, lactone form

 The crude product from Step 2 was dissolved in thioanisole (125 mL), there was added TFA (200 mL) and the mixture was stirred at r.t. for 18 hr. After cooling, the TFA was evaporated, the residue was diluted with Et₂O (400 mL) and after 20 min., the insoluble solid was filtered to afford the desired title compound.

Lactone 2: 3-Hydroxymethyl-4-(3-pyridyl)-7-hydroxy-2-naphthoic acid, lactone form

15



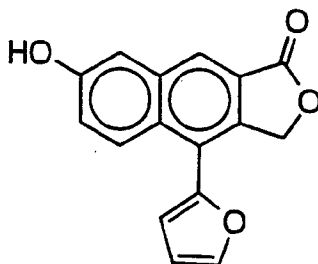
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 Following the procedure described in Lactone 1, Steps 1-3, but substituting 3-pyridinecarbox-aldehyde for 3-furaldehyde, and adding methanesulfonic acid to the trifluoroacetic acid in Step 3, the title compound was obtained.

25

Lactone 3: 3-Hydroxymethyl-4-(2-furyl)-7-hydroxy-2-naphthoic acid, lactone form

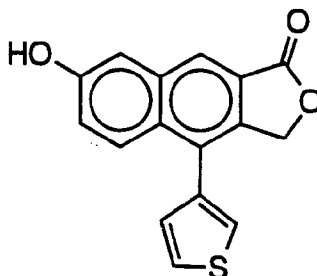
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Following the procedure described in Lactone 1, Steps 1-3, but substituting 2-furaldehyde for 3-furaldehyde, the title compound was obtained.

5 Lactone 4: 3-Hydroxymethyl-4-(3-thienyl)-7-hydroxy-2-naphthoic acid, lactone form



15 Following the procedure described in Lactone 1, Steps 1-3, but substituting 3-thiophenecarboxaldehyde for 2-furaldehyde, the title compound was obtained.

20 The invention is further defined by reference to the following examples, which are intended to be illustrative and not limiting. All temperatures are in degrees Celsius.

EXAMPLE 1

25 3-Hydroxymethyl-4-(3-furyl)-7-[3-[4-(4-hydroxy)tetrahydro-pyranyl]benzyloxy]-2-naphthoic acid, lactone form

30 A mixture of 3-[4-(4-hydroxy)tetrahydropyranyl]benzyl bromide (Halide 2) (355 mg), 3-hydroxymethyl-4-(3-furyl)-7-hydroxy-2-naphthoic acid, lactone form (Lactone 1) (333 mg), and Cs₂CO₃ (426 mg) in DMF (10 mL) was stirred at r.t. overnight. To the reaction mixture was added H₂O (2 mL) followed by EtOAc (100 mL) and the organic layer separated. The aqueous layer was extracted with EtOAc and the combined organic layers were washed with H₂O, brine, dried over MgSO₄ and evaporated. The residue was flash chromatographed on silica gel eluting with hexane:EtOAc:CH₂Cl₂ (1:1:1) to afford the title compound as a white solid; m.p. 181-183°C.

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EXAMPLE 2

5 3-Hydroxymethyl-4-(3-furyl)-7-[3-[4-(4-methoxy)tetrahydro-
pyranyl]benzyloxy]-2-naphthoic acid, lactone form

Following the procedure described in Example 1, but substituting 3-[4-(4-methoxy)tetrahydropyranyl]benzyl bromide (Halide 1) for 3-[4-(4-hydroxy)tetrahydropyranyl]benzyl bromide (Halide 2), the title compound was obtained as a solid; m.p. 179-181°C.

10

EXAMPLE 3

3-Hydroxymethyl-4-(3-pyridyl)-7-[3-[4-(4-hydroxy)-tetrahydro-
15 pyranyl]benzyloxy]-2-naphthoic acid, lactone form

Following the procedure described in Example 1, but substituting 3-hydroxymethyl-4-(3-pyridyl)-7-hydroxy-2-naphthoic acid, lactone form (Lactone 2), for 3-hydroxy-methyl-4-(3-furyl)-7-hydroxy-2-naphthoic acid (Lactone 1), lactone form, the title compound was obtained as a yellow solid; m.p. 219-222°C.

20

EXAMPLE 4

3-Hydroxymethyl-4-(2-furyl)-7-[3-[4-(4-hydroxy)tetrahydro-
25 pyranyl]benzyloxy]-2-naphthoic acid, lactone form

Following the procedure described in Example 1, but substituting 3-hydroxymethyl-4-(2-furyl)-7-hydroxy-2-naphthoic acid, lactone form (Lactone 3), for 3-hydroxymethyl-4-(3-furyl)-7-hydroxy-2-naphthoic acid, lactone form (Lactone 1), the title compound was obtained as a solid; m.p. 164-166°C.

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EXAMPLE 5

3-Hydroxymethyl-4-(3-furyl)-7-[3-[3-(3 α -hydroxy-8-oxabicyclo-
[3.2.1]oct-6-enyl)]benzyloxy]-2-naphthoic acid, lactone form

Following the procedure described in Example 1, but substituting 3-[3-(3 α -hydroxy-8-oxabicyclo-[3.2.1]oct-6-enyl)]benzyl chloride (Halide 3) for 3-[4-(4-methoxy)tetrahydropyranyl]benzyl bromide (Halide 1), the title compound was obtained as a solid; m.p. 222-223°C.

EXAMPLE 6

3-Hydroxymethyl-4-(3-furyl)-7-[3-[3-(3 α -hydroxy-8-oxabicyclo-
[3.2.1]octanyl)]benzyloxy]-2-naphthoic acid, lactone form

Following the procedure described in Example 1, but substituting 3-[3-(3 α -hydroxy-8-oxabicyclo-[3.2.1]octanyl)]benzyl chloride (Halide 4) for 3-[4-(4-methoxy)tetrahydropyranyl]benzyl bromide (Halide 1), the title compound was obtained as a solid; m.p. 195-200°C.

EXAMPLE 7

3-Hydroxymethyl-4-(3-furyl)-7-[5-[4-(4-hydroxy)tetrahydro-
pyranyl]pyridin-3-ylmethoxy]-2-naphthoic acid, lactone form

To a mixture of 3-hydroxymethyl-4-(3-furyl)-7-hydroxy-2-naphthoic acid, lactone form (Lactone 1) (293 mg), 5-[4-(4-hydroxy)tetrahydropyranyl]pyridin-3-ylmethanol (Alcohol 1) (209 mg), and triphenylphosphine (315 mg) in THF (10 mL), there was added di-tert-butyl azodicarboxylate (276 mg) and the mixture was stirred at r.t. for 2 hr. After evaporation of the THF, the residue was flash chromatographed on silica gel, eluting with EtOAc. The product obtained was triturated with ether affording on filtration the title compound as a beige solid; m.p. 122°C (dec).

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EXAMPLE 8

(1S,5R) 3-Hydroxymethyl-4-(3-furyl)-7-[3-[3-(3 α -hydroxy-6,8-dioxabicyclo[3.2.1]octanyl)]benzyloxy]-2-naphthoic acid, lactone form

5 Following the procedure described in Example 7, but substituting (1S,5R) 3-[3-(3 α -hydroxy-6,8-di-oxabicyclo[3.2.1]-octanyl)]-benzyl alcohol (Alcohol 3) for 5-[4-(4-hydroxy)tetrahydropyranyl]-pyridin-3-yl-methanol (Alcohol 1), the title compound was obtained as a solid; m.p. 223°C.

EXAMPLE 9

3-Hydroxymethyl-4-(3-thienyl)-7-[3-[4-(4-hydroxy)-tetrahydropyranyl]benzyloxy]-2-naphthoic acid, lactone form

15 Following the procedure described in Example 1, but substituting 3-hydroxymethyl-4-(3-thienyl)-7-hydroxy-2-naphthoic acid, lactone form (Lactone 4), for 3-hydroxymethyl-4-(3-furyl)-7-hydroxy-2-naphthoic acid, lactone form (Lactone 1), the title compound was obtained as a solid; m.p. 172-175°C.

EXAMPLE 10

3-Hydroxymethyl-4-(3-thienyl)-7-[5-[4-(4-hydroxy)-tetrahydropyranyl]pyridin-3-ylmethoxy]-2-naphthoic acid, lactone form

25 Following the procedure described in Example 7, but substituting 3-hydroxymethyl-4-(3-thienyl)-7-hydroxy-2-naphthoic acid, lactone form (Lactone 4), for 3-hydroxymethyl-4-(3-furyl)-7-hydroxy-2-naphthoic acid, lactone form (Lactone 1), the title compound was obtained as a white solid.

30 ¹H NMR (250 MHz, CDCl₃) δ 1.74 (m, 2H), 1.84 (s, 1H, OH), 2.23 (m, 2H), 3.95 (m, 4H), 5.26 (s, 2H), 5.32 (s, 2H), 7.20-7.60 (m, 5H), 7.90 (d, 1H), 8.00 (s, 1H), 8.37 (s, 1H), 8.68 (s, 1H), 8.78 (s, 1H).

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EXAMPLE 11

(1S,5R) 3-Hydroxymethyl-4-(3-thienyl)-7-[3-(3 α -hydroxy-6,8-dioxabicyclo[3.2.1]octanyl)]benzyloxy]-2-naphthoic acid, lactone form

5 Following the procedure described in Example 7, but substituting (1S,5R) 3-[3-(3 α -hydroxy-6,8-dioxabicyclo[3.2.1]octanyl)]-benzyl alcohol (Alcohol 3) for 5-[4-(4-hydroxy)tetrahydropyranyl]-pyridin-3-ylmethanol (Alcohol 1), and substituting 3-hydroxymethyl-4-(3-thienyl)-7-hydroxy-2-naphthoic acid, lactone form
10 (Lactone 4), for 3-hydroxymethyl-4-(3-furyl)-7-hydroxy-2-naphthoic acid, lactone form (Lactone 1), the title compound was obtained as a solid; m.p. 221-222°C.

EXAMPLE 12

15

(1S,5R) 3-Hydroxymethyl-4-(3-thienyl)-7-[3-[3-(3 α -methoxy-6,8-dioxabicyclo[3.2.1]octanyl)]benzyloxy]-2-naphthoic acid, lactone form

To a solution of (1S,5R) 3-hydroxymethyl-4-(3-thienyl)-7-[3-[3-(3 α -hydroxy-6,8-dioxabicyclo-[3.2.1]octanyl)]benzyloxy]-2-
20 naphthoic acid, lactone form (Example 11) (200 mg) in dry THF (3 mL) at 0°C was added KH (35% in mineral oil, 137 mg, 3 eq.) in dry THF (2 mL). After 10 min. methyl iodide (0.1 mL, 4 eq.) was added and the resulting mixture was stirred for 30 min. at 0°C. The reaction
25 mixture was then poured into saturated aqueous NH₄Cl and extracted with EtOAc. The organic layer was washed with brine, dried over MgSO₄ and evaporated. Flash chromatography on silica gel eluting with EtOAc:hexane (3:7) afforded the title compound as a solid; m.p. 183-184°C.

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EXAMPLE 13

(1R,5S) 3-Hydroxymethyl-4-(3-furyl)-7-[3-[3-(3 α -hydroxy-6,8-dioxabicyclo[3.2.1]octanyl)]benzyloxy]-2-naphthoic acid, lactone form

5 Following the procedure described in Example 7, but substituting (1R,5S) 3-[3-(3 α -hydroxy-6,8-dioxabicyclo[3.2.1]octanyl)]benzyl alcohol (Alcohol 4) for 5-[4-(4-hydroxy)tetrahydropyranol]pyridin-3-ylmethanol (Alcohol 1), the title compound was obtained as a foam.

10 ¹H NMR (300 MHz, CDCl₃): δ 2.00-2.12 (m, 2H), 2.30 (d, 1H), 2.48 (dd, 1H), 3.80 (m, 1H), 3.91 (s, 1H), 4.50 (d, 1H), 4.71 (m, 1H), 5.22 (s, 2H), 5.33 (s, 2H), 5.78 (s, 1H), 6.60 (dd, 1H), 7.19 (m, 1H), 7.37-7.40 (m, 4H), 7.65 (m, 3H), 8.01 (d, 1H), 8.33 (s, 1H).

15

EXAMPLE 14

3-Hydroxymethyl-4-(3-thienyl)-7-[6-[4-(4-hydroxy)-tetrahydropyranyl]pyridin-2-ylmethoxy]-2-naphthoic acid, lactone form

20 Following the procedure described in Example 7, but substituting 6-[4-(4-hydroxy)tetrahydropyran-yl]pyridin-2-ylmethanol (Alcohol 2) for 5-[4-(4-hydroxy)tetrahydropyranyl]pyridin-3-ylmethanol (Alcohol 1), and substituting 3-hydroxymethyl-4-(3-thienyl)-7-hydroxy-2-naphthoic acid, lactone form (Lactone 4), for 3-hydroxymethyl-4-(3-furyl)-7-hydroxy-2-naphthoic acid, lactone form (Lactone 1), the title compound was obtained as a solid.

25 ¹H NMR (250 MHz, CDCl₃): δ 1.6 (m, 2H), 2.15 (m, 2H), 4.0 (m, 4H), 5.15 (s, 1H), 5.3 (s, 2H), 5.35 (s, 2H), 7.2 (dd, 1H), 7.35 (m, 4H), 7.5 (d, 1H), 7.55 (m, 1H), 7.8 (dd, 1H), 7.9 (d, 1H), 8.35 (s, 1H).

30

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EXAMPLE 15

3-Hydroxymethyl-4-(3-furyl)-7-[3-[4-(4-ethyl)tetrahydropyranyl]-benzyloxy]2-naphthoic acid, lactone form

5 Following the procedure described in Example 1, but substituting 3-[4-(4-ethyl)tetrahydropyranyl]-benzyl bromide (Halide 6) for 3-[4-(4-hydroxy)tetrahydropyranyl]benzyl bromide (Halide 2), the title compound was obtained as a pale yellow solid.
10 ¹H NMR (250 MHz, CDCl₃): δ 0.55 (t, 3H), 1.65 (q, 2H), 1.85 (m, 2H), 2.15 (m, 2H), 3.55 (m, 2H), 3.80 (m, 2H), 5.23 (s, 2H), 5.37 (s, 2H), 6.60 (dd, 1H), 7.30-7.45 (m, 6H), 7.65 (m, 2H), 8.02 (d, 1H), 8.33 (s, 1H).

EXAMPLE 16

15

3-Hydroxymethyl-4-(3-furyl)-7-[3-[3-(3 α -hydroxy-2-methoxy-8-oxabicyclo[3.2.1]octanyl)]benzyloxy]-2-naphthoic acid, lactone form

 Following the procedure described in Example 1, but substituting 3-[3-(3 α -hydroxy-2-methoxy-8-oxabicyclo[3.2.1]octanyl)]benzyl chloride (Halide 7) for 3-[4-(4-hydroxy)tetrahydropyranyl]benzyl bromide (Halide 2), the title compound was obtained as a solid; m.p. 185°C.

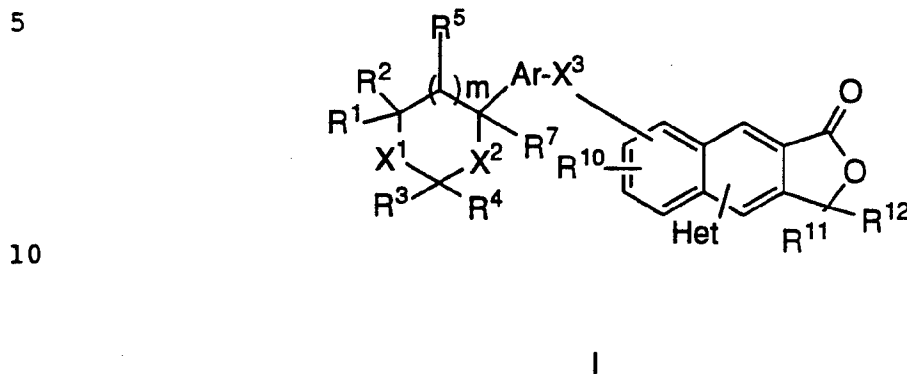
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WHAT IS CLAIMED IS:

1. A compound of the formula:



wherein:

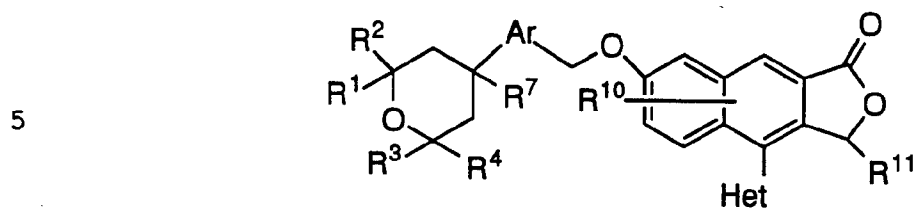
- 15
- R1, R5, and R11 is each independently H, OH, lower alkyl, or lower alkoxy;
- R2 is H, lower alkyl, or together with R1 forms a double bonded oxygen (=O);
- 20 R3 is H, lower alkyl, hydroxy lower alkyl, lower alkoxy lower alkyl, or is joined to R1 to form a carbon bridge of 2 or 3 carbon atoms or a mono-oxa carbon bridge of 1 or 2 carbon atoms, said bridge optionally containing a double bond;
- 25 R4, R12, and R14 is each independently H or lower alkyl;
- R6 is H or lower alkyl, or two R6 groups on the same or adjacent carbons can form a saturated ring of 3 to 8 members;
- R7 is H, OH, lower alkyl, lower alkoxy, lower alkylthio, or lower alkylcarbonyloxy;
- 30 R8 is H, halogen, lower alkyl, lower alkoxy, CF₃, CN, COR¹⁴, or a non-bonded electron pair;
- R9 and R10 is each independently H, lower alkyl, lower alkoxy, hydroxy lower alkyl, lower alkoxy lower alkyl, lower alkylthio lower alkyl, lower alkylthio lower alkylcarbonyl, (R⁸)₂-phenylthio lower alkyl, halogen, CN,

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- NO₂, CF₃, N₃, N(R¹³)₂, NR¹³COR¹⁴, NR¹³CON(R¹³)₂, SR¹⁵, S(O)R¹⁵, S(O)₂R¹⁵, S(O)₂N(R¹³)₂, COR¹⁴, CON(R¹³)₂, CO₂R¹⁴, C(R¹⁴)₂OC(R¹⁴)₂-CO₂R¹⁴, or C(R¹⁴)₂CN;
- 5 R¹⁰ is attached to either ring of the naphtholene ring system;
 R¹³ is H or lower alkyl, or two R¹³ groups attached to the same nitrogen may form a saturated ring of 5 or 6 members, optionally containing a second heteroatom chosen from O, S, or NR⁴;
- 10 R¹⁵ is lower alkyl, phenyl-(R⁸)₂, or CF₃;
 X¹ is O, S, S(O), S(O)₂, or C(R⁶)₂;
 X² is O, S, C(R⁶)₂ or bond;
 X³ is C(R⁶)₂S, SC(R⁶)₂, C(R⁶)₂O, OC(R⁶)₂, CR⁶=CR⁶, C(R⁶)₂C(R⁶)₂, O, or S;
- 15 Ar is arylene(R⁸)₂ wherein arylene is a 5-membered aromatic ring containing one O or S and 0-2 nitrogen atoms; a 5-membered aromatic ring containing 1-3 nitrogen atoms; a 6-membered aromatic ring containing 0-3 nitrogen atoms; 2- or 4-pyranone; or 2- or 4-pyridinone;
- 20 Het is heteroaryl(R⁹)₂ wherein heteroaryl is a 5-membered aromatic ring containing one O or S and 0-3 nitrogen atoms; a 5-membered aromatic ring containing 1-4 nitrogen atoms; a 6-membered aromatic ring containing 1-3 nitrogen atoms; 2- or 4-pyranone; 2- or 4-pyridinone; a bicyclic 8-, 9-, or 10-membered aromatic ring containing 1-4 heteroatoms chosen independently from O, S, and N.
- 25 Het is attached to either ring of the naphthalene ring system; and
 m is 0 or 1; or a pharmaceutically acceptable salt thereof.
- 30

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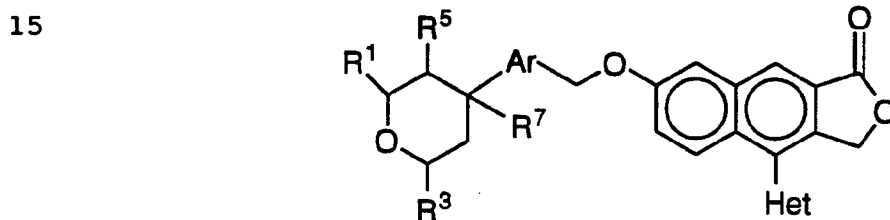
2. A compound of Claim 1 of the formula:



10

or a pharmaceutically acceptable salt thereof.

3. A compound of Claim 2 of the formula:



20

wherein:

25 R¹ is H, Me, or OMe;

R³ is H or Me;

R¹ and R³ together are -CH=CH-, -CH₂CH₂-, -OCH₂-, or -CH₂O-;

R⁵ is H or OCH₃;

30 R⁷ is OH, OCH₃, or CH₂CH₃;

Ar is Phe; 5,3-Pye; 6,2-Pye; 2,4-Pye; or 4,2-Pye; and

Het is 3-Fu or 3-Th;

or a pharmaceutically acceptable salt thereof.

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4. A compound of Claim 3 wherein the substituents are as follows:

EX.	R ¹	R ³	R ⁵	R ⁷	Ar	Het
1	H	H	H	OH	Phe	3-Fu
2	H	H	H	OMe	Phe	3-Fu
3	H	H	H	OH	Phe	3-Py
4	H	H	H	OH	Phe	3-Fu
5	-CH=CH-		H	OH	Phe	3-Fu
6	-CH ₂ CH ₂ -		H	OH	Phe	3-Fu
7	H	H	H	OH	5,3-Pye	3-Fu
8	-CH ₂ O-		H	OH	Phe	3-Fu
9	H	H	H	OH	Phe	3-Th
10	H	H	H	OH	5,3-Pye	3-Th
11	-CH ₂ O-		H	OH	Phe	3-Th
12	-CH ₂ O-		H	OMe	Phe	3-Th
13	-OCH ₂ -		H	OH	Phe	3-Fu
14	H	H	H	OH	6,2-Pye	3-Th
15	H	H	H	Et	Phe	3-Fu
16	-CH ₂ CH ₂ -		OMe	OH	Phe	3-Fu
17	H	H	H	OH	6,2-Pye	3-Fu
18	H	H	H	OH	2,4-Pye	3-Fu
19	H	H	H	OH	4,2-Pye	3-Fu
20	-CH ₂ O-		H	OH	6,2-Pye	3-Fu
21	-CH ₂ O-		H	OH	2,4-Pye	3-Fu
22	-CH ₂ O-		H	OH	4,2-Pye	3-Fu
23	-CH ₂ O-		H	OH	6,2-Pye	3-Fu

5. A pharmaceutical composition comprising a therapeutically effective amount of a compound of Claim 1 and a pharmaceutically acceptable carrier.

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6. A pharmaceutical composition of Claim 5 additionally comprising an effective amount of a second active ingredient selected from the group consisting of non-steroidal anti-inflammatory drugs; peripheral analgesic agents; cyclooxygenase inhibitors; leukotriene antagonists; leukotriene biosynthesis inhibitors; H₁- or H₂-receptor antagonists; antihistaminic agents; prostaglandin antagonists; thromboxane antagonists; thromboxane synthetase inhibitors; and ACE antagonists.
7. A pharmaceutical composition of Claim 6, wherein the second active ingredient is a non-steroidal anti-inflammatory drug.
8. A pharmaceutical composition comprising a therapeutically effective amount of a compound of Claim 1, an effective amount of a second active ingredient which is a non-steroidal anti-inflammatory drug, and a pharmaceutically acceptable carrier, wherein the weight ratio of said compound of Claim 1 to said second active ingredient ranges from about 1000:1 to 1:1000.
9. A method of preventing the synthesis, the action, or the release of SRS-A or leukotrienes in a mammal which comprises administering to said mammal an effective amount of a compound of Claim 1.
10. A method of Claim 9 wherein the mammal is man.
11. A method of treating asthma in a mammal comprising administering to a mammal in need of such treatment a therapeutically effective amount of a compound of Claim 1.
12. A method of treating inflammatory diseases of the eye in a mammal which comprises administering to a mammal in need of such treatment a therapeutically effective amount of a compound of Claim 1.

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13. The method of claim 12 wherein the mammal is man.

14. A leukotriene biosynthesis inhibitor pharmaceutical composition comprising an acceptable leukotriene biosynthesis
5 inhibiting amount of a compound or salt as defined in Claim 1, 2, 3 or 4, in association with a pharmaceutically acceptable carrier.

15. A compound or salt, as defined in Claim 1, 2, 3 or 4 for use in the treatment of asthma or inflammatory diseases.

16. Use of a compound or salt, as defined in claim 1, 2,
10 3 or 4, for the manufacture of a medicament for the treatment of asthma or inflammatory diseases.

INTERNATIONAL SEARCH REPORT

International Application No

PCT/CA 93/00339

A. CLASSIFICATION OF SUBJECT MATTER

IPC 5 C07D407/14 C07D405/14 C07D493/08 C07D409/14 A61K31/34
 A61K31/38 A61K31/44 //(C07D493/08,311:00,307:00),
 (C07D493/08,317:00,311:00)

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 5 C07D

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	EP,A,0 188 248 (TANABE SEIYAKU CO.,LTD) 23 July 1986 see claims 1-3,8 ---	1,5-16
A	EP,A,0 409 413 (IMPERIAL CHEMICAL INDUSTRIES PLC) 23 January 1991 see claims 1,12 ---	1,5-16
P,X	EP,A,0 501 579 (MERCK FROSST CANADA INC.) 2 September 1992 see claims 15-10 -----	1,5-16

☐ Further documents are listed in the continuation of box C.☒ Patent family members are listed in annex.

* Special categories of cited documents:

"A" document defining the general state of the art which is not considered to be of particular relevance

"E" earlier document but published on or after the international filing date

"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)

"O" document referring to an oral disclosure, use, exhibition or other means

"P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.

"&" document member of the same patent family

Date of the actual completion of the international search

15 November 1993

Date of mailing of the international search report

26. 11. 93

Name and mailing address of the ISA

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Authorized officer

HENRY, J

INTERNATIONAL SEARCH REPORT

International application No.

PCT/CA93/00339

Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)

This international search report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☐ Claims Nos.:
because they relate to subject matter not required to be searched by this Authority, namely:
Although claims 9-13 are directed to a method of treatment of human body, the search has been carried out and based on the alleged effects of the compounds.
2. ☐ Claims Nos.:
because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out, specifically:
3. ☐ Claims Nos.:
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

1. ☐ As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims.
2. ☐ As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3. ☐ As only some of the required additional search fees were timely paid by the applicant, this international search report covers only those claims for which fees were paid, specifically claims Nos.:
4. ☐ No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

Remark on Protest

- ☐ The additional search fees were accompanied by the applicant's protest.
- ☐ No protest accompanied the payment of additional search fees.

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/CA 93/00339

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
EP-A-0188248	23-07-86	AU-B- 584153	18-05-89
		AU-A- 5175185	17-07-86
		JP-A- 61267541	27-11-86
		JP-A- 2072136	12-03-90
		JP-A- 2072170	12-03-90
		US-A- 4897418	30-01-90
		US-A- 5070103	03-12-91
		US-A- 4771072	13-09-88
EP-A-0409413	23-01-91	AU-B- 636357	29-04-93
		AU-A- 5757590	24-01-91
		CA-A- 2019345	18-01-91
		JP-A- 3056474	12-03-91
		US-A- 5208259	04-05-93
EP-A-0501579	02-09-92	JP-A- 5140143	08-06-93
		US-A- 5227399	13-07-93